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OM protein - protein search, using sw model	
Run on:	August 22, 2005, 10:02:08 ; Search time 178 Seconds (without alignments)
Title:	US-10-629-951-2
Perfect score:	3055
Sequence:	1 MGTGLESLSLGDGRGANPTV..... HKLNKVDWLMENHEKUSLS 580
Scoring table:	BLOSUM62
Searched:	Gapop 10.0 , Gapext 0.5
Total number of hits satisfying chosen parameters:	1612378
Minimum DB seq length:	0
Maximum DB seq length:	2000000000
Post-processing:	Minimum Match 0% Maximum Match 100% Listing first 45 summaries
Database :	Uniprot 03: 1: uniprot_sprot: 2: uniprot_trembl: Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.
SUMMARIES	
Result No.	Score
1	3055
2	1821.5
3	880.5
4	344
5	338
6	238.5
7	220.5
8	213
9	208.5
10	208.5
11	208.5
12	207.5
13	202.5
14	201
15	198.5
16	156
17	149.5
18	148.5
19	148.5
20	146
21	146
22	145
23	145
24	145
25	144
26	143.5
27	141.5
28	141.5
29	140.5
30	140.5
31	140.5
32	140.5
33	138.5
34	138.5
35	138.5
36	137.5
37	137.5
38	137.5
39	137.5
40	137
41	137
42	137
43	137
44	137
45	136.5
ALIGNMENTS	
Query Match Length DB ID	Description
1 MBP4_HUMAN	095243 homo sapien
2 MBP4_MOUSE	09z2d7 mus musculus
3 Q919F1	Q919F1 gallus gallus
4 Q8AFCL	Q8AFCL arabidopsis
5 Q84WT3	Q84WT3 arabidopsis
6 Q7T2T7	Q7T2T7 brachydorina
7 Q9YGC6	Q9YGC6 xenopus laevis
8 Q7SCQ2	Q7SCQ2 neurospora crassa
9 MEC2_HUMAN	P51608 homo sapien
10 Q6QHH9	Q6QHH9 homo sapien
11 Q72384	Q72384 homo sapien
12 MEC2_MACPA	Q95198 macaca fasciata
13 Q42403	Q42403 gallus gallus
14 MEC2_RAT	Q00566 rattus norvegicus
15 MEC2_MOUSE	Q922d6 mus musculus
16 Q7RN41	Q7RN41 plasmodium
17 Q811T6	Q811T6 plasmodium
18 Q8LB76	Q8LB76 arabidopsis thaliana
19 Q7RCP2	Q7RCP2 plasmodium
20 Q8SF22	Q8SF22 arabidopsis thaliana
21 Q8HQJ1	Q8HQJ1 pan troglodytes
22 Q8AYT1	Q8AYT1 xenopus laevis
23 P46674	P46674 saccharomyces cerevisiae
24 Q81b76	Q81b76 homo sapien
25 Q8AYP2	Q8AYP2 xenopus laevis
26 Q25875	Q25875 plasmodium
27 Q25730	Q25730 plasmodium
28 Q26007	Q26007 plasmodium
29 Q8U414	Q8U414 plasmodium
30 Q8U430	Q8U430 plasmodium
31 Q7RK82	Q7RK82 plasmodium
32 Q924D9	Q924D9 meriones shawii
33 P41891	P41891 schizosaccharomyces pombe
34 Q9u429	Q9u429 plasmodium
35 Q7RPJ9	Q7RPJ9 plasmodium
36 Q869D5	Q869D5 branchiostoma floridae
37 Q6MF11	Q6MF11 neurospora crassa
38 Q26104	Q26104 plasmodium
39 Q9u0G6	Q9u0G6 plasmodium
40 Q81k06	Q81k06 zea mays (millet)
41 P42348	P42348 baccharomyces
42 Q7RKH5	Q7RKH5 plasmodium
43 Q7t150	Q7t150 mus musculus
44 Q75X83	Q75X83 helicobacter pylori
45 P35663	P35663 homo sapiens

RA Ebert L., Schick M., Neubert P., Schatten R., Henze S., Korn B.;
 RT "Cloning of human full open reading frames in Gateway(TM) system entry
 vector (pDONR011);";
 RT submitted (MAY-2004); to the EMBL/GenBank/DBJ databases.
 RN [7]
 RP SEQUENCE FROM N.A. (ISOFORM 2).
 TISSUE=Lung;
 MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
 RX STRAUSSBERG R.L., Fengold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Sheinmen C.M., Schuler G.D.,
 RA Altshul S.F., Zeeberg B., Buetow K.H., Schaeffer C.F., Bhat N.K.,
 RA Bosak S.A., McBwan P.J., McKernan K.J., Malek J.A., Gunnarino P.H.,
 RA Richard R.P., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diachenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Uddin T.B., Toshiyuki C., Prange J.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzywinski M.T., Stalska U., Smailus D.E.,
 RA Schnarch A., Schein J.E., Jones S.J.M., Marra M.A.,
 RA RT "Generation and initial analysis of more than 15,000 full-length human
 and mouse cDNA sequences";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
 RN [8]
 RP FUNCTION.
 RX Pubmed=0930409; DOI=10.1074/jbc.M0045355200;
 RA Petronzelli F., Riccio A., Markham G.D., Seeholzer S.H., Stoerker J.,
 RA Genardi M., Yeung A.T., Matsumoto Y., Bellacosa A.;
 RT "Biphasic kinetics of the human DNA repair protein MED1 (MBD4), a
 RT mismatch-specific DNA N-glycosylase.";
 RL J. Biol. Chem. 275:32422-32429 (2000).
 RN [9]
 RP INTERACTION WITH FADD.
 RX Pubmed=1270275; DOI=10.1073/pnas.0431215100;
 RA Screenon R.A., Kiessling S., Sansom O.J., Millar C.B., Maddison K.,
 RA Bird A., Clarke A.R., Frisch S.M.;
 RT "Fas-associated death domain protein interacts with methyl-CPG binding
 RT domain protein 4: a potential link between genome surveillance and
 RT apoptosis";
 RL Proc. Natl. Acad. Sci. U.S.A. 100:5211-5216 (2003).
 RT -!- FUNCTION: Mismatch-specific DNA N-glycosylase involved in DNA
 repair. Has thymine glycosylase activity and is specific for G:T
 mismatches within methylated and unmethylated Cpg sites. Can also
 remove uracil or 5-fluorouracil in G:U mismatches. Has no lyase
 activity. Was first identified as methyl-CpG-binding protein.
 CC --!- SUBUNIT: Interacts with MBD1.
 CC --!- SUBCELLULAR LOCATION: Nuclear.
 CC --!- ALTERNATIVE PRODUCTS:
 Event=Alternative splicing; Named isoforms=3;
 Note=No experimental confirmation available;
 CC ISOId=095243-1; Sequence=Displayed;
 Name=2;
 ISOId=095243-2; Sequence=VSP 010816;
 Note=No experimental confirmation available;
 Name=3;
 ISOId=095243-3; Sequence=VSP 010817, VSP 010818;
 Note=No experimental confirmation available;
 CC --!- SIMILARITY: Contains 1 methyl-CpG-binding (MBD) domain.

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 CC DR EMBL; AF072250; AAC68879.1; -.

DR EMBL; AF120999; AA050374.1; -.
 DR EMBL; AF120997; AA050374.1; JOINED.
 DR EMBL; AF120998; AA050374.1; -.
 DR EMBL; AF114784; AA022195.1; -.
 DR EMBL; AF532602; RA977338.1; -.
 DR EMBL; AF494057; AA000088.1; -.
 DR EMBL; CR450305; CGG29301.1; -.
 DR EMBL; BC011752; AAH11752.1; -.
 DR HSSP; Q922D7; INGN.
 DR InAct; O95243; -.
 DR Genew; HGNC6919; MBD4.
 DR H-InvDB; HIK0031659; -.
 DR Reactome; O95243; -.
 DR MM; 603574; -.
 DR GO; GO:0005634; C-nucleus; TAS.
 DR GO; GO:0004520; Endo-ribonuclease activity; TAS.
 DR GO; GO:0003636; F-satellite DNA binding; TAS.
 DR GO; GO:0006281; RNA repair; TAS.
 DR InterPro; IPR003205; Endo_3c.
 DR InterPro; IPR001739; Methy-CpG_bind.
 DR Pfam; PF00730; HmN-GPD; 1.
 DR Pfam; PF01428; MBD; 1.
 DR SMART; SM00321; MBD; 1.
 DR PROSITE; PS50982; MBD; 1.
 KW Alternative splicing; DNA repair; DNA-binding; Hydrolase;
 KW Nuclear protein; Polymerase.
 FT DOMAIN 76 148 MBD.
 FT DOMAIN 461 524 HmN-GPD.
 FT ACT_SITE 560 560 BY similarity.
 FT VARSPLIC 395 400 Missing (in Isoform 2).
 FT VARSPLIC 539 540 /FTid=VAR 010816.
 FT VARSPLIC 541 580 KY -> AP (In Isoform 3).
 FT VARSPLIC 541 580 /FTid=VAR 010817.
 FT VARIANT 342 342 S -> P (In dbsnp:2307289).
 FT VARIANT 346 346 /FTid=VAR 019358.
 FT VARIANT 358 358 /FTid=VAR 019514.
 FT VARIANT 568 568 E -> K (In dbsnp:140693).
 FT VARIANT 580 580 /FTid=VAR 019359.
 FT SEQUENCE 580 AA: BPI6PP2IA3ABE5F CRC64;
 QY 1 MGTTGLEISLSLGDRGAAPTVTSSRLVDPNDPLRKEDVAMELERVGDEEQNMKRSSE 60
 Db 1 MTGTGLEISLSLGDRGAAPTVTSSRLVDPNDPLRKEDVAMELERVGDEEQNMKRSSE 60
 Db 121 KSSLANYIHKNGETSLKPDPDFVLSRGKISRYKDCSMAALTSHLQNSNNWIRT 180
 Db 121 KSSLANYIHKNGETSLKPDPDFVLSRGKISRYKDCSMAALTSHLQNSNNWIRT 180
 QY 61 CNPQIQLQEPFLASAORGATAGTCERKSVPGWERTVKQRFGKTAGFDTYFISQGKRS 120
 Db 61 CNPQIQLQEPFLASAORGATAGTCERKSVPGWERTVKQRFGKTAGFDTYFISQGKRS 120
 QY 121 KSSLANYIHKNGETSLKPDPDFVLSRGKISRYKDCSMAALTSHLQNSNNWIRT 180
 Db 181 RSKCKDVFMPSSSETQESRGISNFSTHLLIKEDGVDDYNFRKTKPKRKVTLLG 240
 QY 241 IPIKTKKGCRKCSGFGFQDSKRESVONKADESEPAQKSOLDRTVCISAGCETL 300
 Db 241 IPIKTKKGCRKCSGFGFQDSKRESVONKADESEPAQKSOLDRTVCISAGCETL 300

RESULT 3

Q91P1 PRELIMINARY; PRT; 416 AA.

AC Q91P1;

DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)

DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)

DE 5-methylcytosine G/T mismatch-specific DNA glycosylase.

OS Gallus gallus (Chicken)

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Buteleostomi; Archosauria; Aves; Neognathae; Galiformes; Phasianidae; Phasianinae; Gallus.

OX NCBI_TaxID=9031;

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE=0512662; PubMed=1108112; DOI=10.1093/nar/28.21.4157;

RA Zhiu B., Zheng Y., Angliker H., Schwarz S., Thiry S., Siegmann M., Jost J.-P.,

RT "5-Methylcytosine DNA glycosylase activity is also present in the human MBD4 (G/T mismatch glycosylase) and in a related avian sequence," Nucleic Acids Res. 28:4157-4165 (2000).

DR EMBL; AF251707; AAF68981.1; -.

DR HSSP; Q9ZD7; INGN.

DR InterPro; IPR011257; DNA_glycosyle.

SQ SEQUENCE: 416 AA; 45454 MW; A3F70C6FF2133F2A CRC64;

Query Match 28.8%; Score 880.5; DB 2; Length 416; Best Local Similarity 78.2%; Pred. No. 1.2e-45; Matches 161; Conservative 20; Mismatches 21; Indels 4; Gaps 2;

QY 378 RGSEMDNNCS--PTRKDFGKIFQ--EDTPRTQIERRKLISLYFSSSKYKNEALKSPPRK 433

Db 210 RDSEAAGDVSPSPDKGSFTAVQAPRGTEESAPRTQYDRRKPSVSSKYSKEALKSPPRK 269

QY 434 AFKKWTPPRSPFPNLVQBTLEFDPWKLIAITFLNKTSKMAIPVLFIRKYPSEAVRT 493

Db 270 AFRKKWTPPRSPFPNLVQBTLEFDPWKLIAITFLNKTSKMAIPVLFIRKYPSEAVRT 329

QY 494 ADWRYVSELRLPLGLYDLRAKTIVFSDETLTQKQYPIHLHGIGKYGDSYRIFCVNEW 553

Db 330 ADWKENSELLRPLGLYALRATKIKFSEDEVINKQKPYIELHGIGKYGDSYRIFCVNEW 389

QY 554 KQVHEDHKUNQHYHDLWENHEKLSL 579

Db 390 KEVQPOGDHKUNIYHAWLWENHEKLSV 415

RESULT 4

Q9SFC1 PRELIMINARY; PRT; 419 AA.

AC Q9SFC1;

DT 01-MAY-2000 (TREMBLrel. 13, Created)

DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)

DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)

DR Hypothetical protein At3g07930.

GN Name=At3g07930;

OS Arabidopsis thaliana (Mouse-ear cress)

OC Eukaryota; Viridiplantae; Striphophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

OX NCBI_TaxID=3702;

RN [1]

RP SEQUENCE FROM N.A.

RA Yamada K., Chan M.M., Chang C.H., Dale J.M., Huan V.W., Lee J.M., Onodera C.S., Quach H.L., Tang C., Toriumi M., Wong C., Wu H.C., Yu G., Yuan S., Carninci P., Chen H., Cheuk R., Hayashizaki Y., Ishida J., Jia T., Kamiya A., Kawai J., Kim C.J., Narasaki M., Nguyen M., Palm C.J., Sakurai T., Sakou M., Seki M., Shim P., Southwick A., Tripp M.G., Wu T., Shinozaki K., Davis R.W., Ecker J.R., Arapidopsis thaliana (Mouse-ear cress).

RESULT 5

Q84WT3 PRELIMINARY; PRT; 407 AA.

AC Q84WT3;

DT 01-JUN-2003 (TREMBLrel. 24, Created)

DT 01-JUN-2003 (TREMBLrel. 24, Last sequence update)

DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)

DR Hypothetical protein At3g07930.

GN Name=At3g07930;

OS Arabidopsis thaliana (Mouse-ear cress)

OC Eukaryota; Viridiplantae; Striphophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

OX NCBI_TaxID=3702;

RN [1]

RP SEQUENCE FROM N.A.

RA Yamada K., Chan M.M., Chang C.H., Dale J.M., Huan V.W., Lee J.M., Onodera C.S., Quach H.L., Tang C., Toriumi M., Wong C., Wu H.C., Yu G., Yuan S., Carninci P., Chen H., Cheuk R., Hayashizaki Y., Ishida J., Jia T., Kamiya A., Kawai J., Kim C.J., Narasaki M., Nguyen M., Palm C.J., Sakurai T., Sakou M., Seki M., Shim P., Southwick A., Tripp M.G., Wu T., Shinozaki K., Davis R.W., Ecker J.R.,

RA	Theologis A.;	Matches 106; Conservative 57; Mismatches 159; Indels 113; Gaps 18;
RL	Submitted (JAN-2003) to the EMBL/GenBank/DBBJ databases.	
DR	EMBL; BT002799; AA02623.1; -.	
DR	HSSP; Q9ZD7; 1NGN.	
DR	GO; GO:0006284; P:base-excision repair; IEA.	
DR	InterPro; IPR011257; DNA_glycosylase.	
DR	InterPro; IPR003265; Endo_3c.	
DR	Pfam; PF00730; HhH_GPD; 1.	
KW	Hypothetical protein.	
SQ	SEQUENCE 407 AA; 46941 MW; DD758CD862EFS4F CRC64;	
	Query Match 11.1%; Score 338; DB 2; Length 407; Best Local Similarity 26.6%; Pred. No. 1. 4e-12; Mismatches 155; Indels 76; Gaps 11; Matches 102; Conservative 51; Mismatches 155; Indels 76; Gaps 11;	
Qy	251 RKCSGSPVQ-SDSKRESCVNKADESPPVACKSQUALRTVCISDAGGETTSVTSBENSL 309	
Db	30 RRPDSDFIEVDENRSPALPKEDDEK-----NRDLGIYDDGSTNLVHQCHONGCL 80	
Qy	310 VKKERKSLS--SGSNICSEOK-----TSGINKFRSAKOSHEHNEXCVDITLESE 356	
Db	81 EKDONSISLDDPLFSGFVTKGVRRKRDKDFGSGTITSNLVSPQADD---DDVSVDHSIERO 137	
Qy	357 EIGTKVKEVERK-----ERLHTDILKRGSEMID-NCSPTRKPFGEK--FOEDT 403	
Db	138 ECISKVQAKPVRSVPYFOASTISQCDSDIVSSQSGNSQRKSYRKSSKROVKARRSPVTFQEST 197	
Qy	404 IP-----RTQLERKUSLYF-----SSKYNKRALSP----- 430	
Db	198 VSEQPNOAPKGGLANRYFKVVKSYRFHDGIONESOREKSKRNVRKTPIVSPVLSQKTD 257	
Qy	431 ---RRAKAFKKTIPRSPFNLVOETLFLHDPMKLLIATIFURTSGMIALPIWLKFELKYP 486	
Db	258 DYLRLKPTPDNTWVPPRECNPNCQEDHMPWVPLVTCMLNLTSGAQTRGVISDLRFLCT 317	
Qy	318 SAVBARTADWODSELKPLGLAKTIVKRESDEVLTTRKQYKTFELHGKIGNDIR 546	
Db	547 IFCVNEWKQVHEDHKUNKYHML 570	
Db	378 IFCGNNDRVKENDHMLNYWVDYL 401	
	RESULT 6	
ID	07T2T7 PRELIMINARY; PRT; 524 AA.	
AC	07T2T7; 2003 (TREMBrel. 25, Created)	
DT	01-OCT-2003 (TREMBrel. 25, Last sequence update)	
DT	01-MAR-2004 (TREMBrel. 26, Last annotation update)	
DE	Methyl-L-cysteine binding protein 2.	
GN	Name=mecp2;	
OS	Brachydanio rerio (zebrafish) (Danio rerio).	
OC	Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
OC	Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;	
OC	Cyprinidae; Danio.	
OX	NCBI_TaxId=7955;	
[1]	SEQUENCE FROM N.A.	
RA	Coverdale L.B., Martin C.C.;	
RL	Submitted (Oct-2003) to the EMBL/GenBank/DBBJ databases.	
DR	EMBL; AA02624.2; -.	
EMBL; AU298300; AAP57248.2; -.		
DR	HSSP; Q9UNZ9; 1IG4.	
DR	ZFIN; ZDB-GENE 030131-7190; mecp2.	
DR	GO; GO:0003677; F:DNA binding; IEA.	
DR	InterPro; IPR001739; Methylation_CpG_bind.	
DR	Pfam; PF01429; MBD; 1.	
DR	SMART; SM00331; MBD; 1.	
SQ	SEQUENCE 524 AA; 57152 MW; B8593B4BB84DC2D1 CRC64;	
	Query Match 7.8%; Score 230.5; DB 2; Length 524; Best Local Similarity 24.4%; Pred. No. 2. 2e-06; Length 524;	
Qy	27 VPDPNDLRKVEDAMELERVEDEQMMIKRSSENCPNLIQEPEITASAQFGA---TAGTECR 83	
DR	49 VPPPSLFTQRDVEGQAE-AGKSE-----PI--DPEVGAAALSAPESSASAKOR 93	
DR	84 KSV-----PCGMRVVKQRLFAGTAGPDRVYFISPOCIKFRSKSSLANYLHK 130	
DR	94 RSVIRDGGPMYEDSLPQGWTRKUKQKGRSACKFDDVULINPEGKAFRSKVELMAYRQK 153	
Db	131 NGETLKLKPERDFVTLSKORGKSRVYKDCSMAALTSHLQMQNSNNWNTRSKCKDVF 190	
Db	154 VGDTITPDNFDFTV-TGRGSPR-----RBKRPKPKRQW 188	
Qy	191 PPSSSELQESRCGLSNFTSHLINKEDEGYDDVNRKVRKPKGKVTI----- 237	
Db	189 KPS-----GRGRGPKGSKGKVROATEGV-AVKRVIBKSPGKLUVKMPFVAPKTEPGA 239	
Qy	238 -LKGIPKCTKKGCRKCSGFVQSDSKRESVCNKADAEPPVAKQSQUALRTVCISDAGAC 296	
Db	240 PLGQAPVAKARRGRK-----KSQDPSTPKRGRKRPAVTSQS----TUGHTSAAY 288	
Qy	297 GETLSYSE-----ENSLVKKERSLSSGSNFCSQEQTSGIINKFCSAKDSEHNEKYE 349	
Db	289 AAAATTAEAKKALKKESAKPVOAKLP-----IKRKRCRETLBL-EASTSATETFE 342	
Qy	350 DTFLESEBIGKVEV-VERCEHLHDILKRGSEMDNNNCSPTRKDFTGEGKIFOEDTRPTQ 408	
Db	343 KRLTASTVTPGEEABTGQPKHPS-RICHKEADPGSSSSGTASG-----VAPKSH 393	
Qy	409 IERRKTSLSYSSKYN 423	
Db	394 KKRDRQQHFKHHH 408	
	RESULT 7	
ID	09YGC6 PRELIMINARY; PRT; 467 AA.	
AC	09YGC6; 1999 (TREMBrel. 10, Created)	
DT	01-MAY-1999 (TREMBrel. 10, Last sequence update)	
DT	01-MAY-1999 (TREMBrel. 10, Last annotation update)	
DE	Methyl-CpG-binding protein 2.	
GN	Name=mecp2;	
OS	Xenopus laevis (African clawed frog).	
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
OC	Amphibia; Batrachia; Anura; Mesobatrachia; Pipoidea; Pipidae;	
OC	Xenopoda; Xenopidae; Xenopus.	
OX	NCBI_TaxId=8355;	
RN	[1]	
RP	SEQUENCE FROM N.A.	
RX	MEDLINE=9882101; PubMed=9620779; DOI=10.1038/561; Jones P.L., Veerstra G.J.C., Wade P.A., Vermaak D., Kass S.U., Landberger N., Strubbe J., Wolffe A.P.; "Methylated DNA and MeCP2 recruit histone deacetylase to repress transcription." Nat. Genet. 19:187-191(1998).	
RT		
RN	[2]	
RP	SEQUENCE FROM N.A.	
RA	Kass S.U., Strubbe J., Wolffe A.P.;	
RL	Submitted (FEB-1998) to the EMBL/GenBank/DBBJ databases.	
DR	EMBL; AF10951; AAD0736.1; -.	
DR	EMBL; AF051768; AAD02651.1; -.	
DR	HSSP; P51608; 1OK3.	
DR	GO; GO:0005634; C:nucleus; IEA.	
DR	GO; GO:003677; F:DNA binding; IEA.	
DR	GO; GO:0006355; P:regulation of transcription, DNA-dependent; IEA.	
DR	InterPro; IPR001737; A+T hook.	
DR	InterPro; IPR001739; Methylation_CpG_bind.	
DR	Pfam; PF02178; AT hook; 1.	
DR	SMART; SM00384; AT hook; 2.	
DR	SMART; SM00391; MBD; 1.	

DR InterPro; IPR003265; Endo_3c.
 DR PFam; PF00730; HhB-GPD; 1.
 DR Hypothetical protein.
 SQ SEQUENCE 682 AA; 4FDCCAA26102E8E4 CRC64;
 Best Local Similarity 20.8%; Pred. No. 0.0001; Length 682;
 Matches 93; Conservative 62; Mismatches 155; Indels 138; Gaps 14;
 DR SEQUENCE 75928 MW; 4FDCCAA26102E8E4 CRC64;
 Best Local Similarity 20.8%; Pred. No. 0.0001; Length 682;
 Matches 93; Conservative 62; Mismatches 155; Indels 138; Gaps 14;
 DR SEQUENCE 7.0%; Score 213; DB 2; Length 682;
 DR SEQUENCE 100 MW; 5D3A719A5E560BC CRC64;
 Best Local Similarity 25.7%; Pred. No. 2.3e-05; Length 467;
 Matches 98; Conservative 60; Mismatches 157; Indels 67; Gaps 15;
 DR SEQUENCE 22 MW; 5D3A719A5E560BC CRC64;
 Best Local Similarity 25.7%; Pred. No. 2.3e-05; Length 467;
 Matches 98; Conservative 60; Mismatches 157; Indels 67; Gaps 15;
 DR SEQUENCE 22 SSERLVPDPNDLRKEDVAMELER--VGSEDEQMIKRS--SECNPLIOPPIASAQFGA 76
 DR SEQUENCE 49 MW; 5D3A719A5E560BC CRC64;
 Best Local Similarity 25.7%; Pred. No. 2.3e-05; Length 467;
 Matches 98; Conservative 60; Mismatches 157; Indels 67; Gaps 15;
 DR SEQUENCE 77 TAGTECRKSVCPCGWERVVKOFGLFGKTAGRDTVYFTSPQGKLRFSKSLANYLHKNGETS 136
 DR SEQUENCE 100 MW; 5D3A719A5E560BC CRC64;
 Best Local Similarity 25.7%; Pred. No. 2.3e-05; Length 467;
 Matches 98; Conservative 60; Mismatches 157; Indels 67; Gaps 15;
 DR SEQUENCE 137 KPEDDFPTVLSKRGTSRYKDCSMLALTSHLONDNSNNWNLTRSKCKDV--FMPSS 194
 DR SEQUENCE 152 DPNDFDFTV-TGGRPSRREQ----KOPKKPKAPKSSVGRGRGRPKCSIKVKKPPV 204
 DR SEQUENCE 195 SSELQSRGLSNFTSHLLIK----EDEGVDDVNPKV----RKEPKGVTLKGIP 242
 DR SEQUENCE 205 SEGVQYQTKVTK-SPGKLVTPMVSQSGTKEASDATTSQSVLVIKGGRKORKSE-TDPSAAP 262
 DR SEQUENCE 243 IKKTKKGCRKSCSGFVQDSKRSKESVNCNADESERVAQKSQLDRTVCISDAAGCETLSV 302
 DR SEQUENCE 263 KKRGRKPSNVLAAAEEAKKKAI--KESKILIE----TVLPIKKRKRETISV 313
 DR SEQUENCE 303 TSEE-----NSLYRK-----KERSLSSGSNFNSEQKTSQGTLINKFCACKSHNEKCE 349
 DR SEQUENCE 314 DVKDTIKPEPITPVIEKVMKQNPANSPEPSTEGSPKLIKGLPKKELQOHHHHHHHH 373
 DR SEQUENCE 350 DTFLESEEIGTKVEVERKEHHL 371
 DR SEQUENCE 374 HHHSBSKASATSPPEPETSQDNII 395

RESULT 8

Q7SCQ2 PRELIMINARY; PRT; 682 AA.

AC Q7SCQ2; 01-MAR-2004 (TREMBl; 26, Created)
 DT 01-MAR-2004 (TREMBl; 26, Last sequence update)

DT 01-MAR-2004 (TREMBl; 26, Last annotation update)
 DR Hypothetical protein.
 Name=NCU09815;1;

GN Neurospora crassa.

OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;

OC Sordariomycetidae; Sordariales; Sordariaceae; Neurospora.

OX NCBI_TAXID=511; [1]

DR SEQUENCE FROM N.A.

DR STRAIN=OR4A;

RA Galagan J.E., Calvo S.E., Borkovich K.A., Selker E.U., Read N.D.,
 Jaffee D., Fitzhugh W., Ma L.-J., Smirnov S., Purcell S., Rehman B.,
 Elkins T., Engels R., Wang S., Nielsen C.B., Butler J., Endrizzi M.,
 Qui D., Janakiev P., Pedersen D., Nelson M., Washburn M.,
 Sellitremnikoff C.P., Kinsey J.A., Braun E.L., Zelter A., Schulte U.,
 Kothe G.O., Jedd G., Meves W., Staben M., Marcotte E., Greenberg D.,
 Roy A., Foley K., Naylor J., Thomann N., Barrett R., Gnerre S.,
 Kamal M., Kamysseis M., Mauceli E., Belke C., Rudd S., Frisman D.,
 Krystofova S., Rasmussen C., Metzenberg R.L., Perkins D., Kroken S.,
 Coconi C., Macino G., Catcheside D., Li W., Pratt R.J., Osmani S.A.,
 DeSouza C., Glass L., Orbach M.J., Berglund J., Voecker R.,
 Yarden O., Plamann M., Seiler S., Dunlap J., Radford A., Aramayo R.,
 Natvig D.O., Alex T.A., Mannhaupt G., Ebbole D.J., Freitag M.,
 Paulsen I., Sachs M.S., Lander E.S., Nusbaum C., Birren B.,
 RT "The Genome Sequence of the Filamentous Fungus Neurospora crassa.";
 RL Nature 0 0-0 (2003).
 CC -!- CAUTION: The sequence shown here is derived from an
 EMBL/GenBank/DDBJ whole genome shotgun (WGS) entry which is
 preliminary data.

DR GO; GO:0006284; P-base-excision repair; IEA.
 DR Intq-Pro; IP011257; DNA_glycosylse.

DR GO; GO:0006284; P-base-excision repair; IEA.

RESULT 9

MEC2_HUMAN STANDARD; PRT; 486 AA.

AC P31608; 015233; 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)

DT 25-OCT-2004 (Rel. 45, Last annotation update)
 DE Methyl-Cpg-binding protein 2 (MeCP-2 protein) (MeCP2).
 Name=MECP2;

OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Catarrini; Hominoidea; Homo.

OX NCBI_TAXID=9606; [1]

DR SEQUENCE FROM N.A.

RA Kudo S., Fukuda M.; Submitted (SBP-1995) to the EMBL/GenBank/DDBJ databases.

RA SEQUENCE FROM N.A.

RA TissuePlacenta; Thiesen J.; Sträatling W.H.; Submitted (APR-1997) to the EMBL/GenBank/DDBJ databases.

RA [3] Submitted (APR-1997) to the EMBL/GenBank/DDBJ databases.

RA SEQUENCE FROM N.A.

RA MEDLINE-97130625; Published=8376388;

RA Vilain A., Apiou F., Vogt N., Dutrillaux B., Malfoy B.; Methyl-Cpg-binding protein 2 (MECP2) to

RT human chromosome band xq28 by *in situ* hybridization."

RT Cyogenet. Cell Genet. 74:293-294(1996).

RN [4] SEQUENCE FROM N.A.

RP SEQUENCE FROM N.A.

- RA Reichwald K., Rosenthal A., Kioschis P., Platzer M.; "Mapping and sequence analysis of the human MeCP2 locus."; submitted (OCT-1997) to the EMBL/GenBank/DDBJ databases.
- [15]
- RN SEQUENCE FROM N.A.
- RP MEDLINE=99299249; PubMed=10369871; DOI=10.1093/hmg/8.7.1253;
- RA Coy J. F., Sedlacek Z., Brechner D., Delius H., Poustka A.;
- RT "A complex pattern of evolutionary conservation and alternative polyadenylation within the long 3'-untranslated region of the methyl-CpG-binding protein 2 gene (MECP2) suggests a regulatory role in gene expression."; Hum. Mol. Genet. 8:1253-1262(1999).
- RL [16]
- RP SEQUENCE FROM N.A.
- RA Reichwald K., Thiesen J., Wiehe T., Weitzel J., Straetling W.H., Kioschis P., Rosenthal A., Platzer M.; "Comparative sequence analysis of the MECP2-locus in human and mouse reveals new untranslated regions."; Submitted (JUN-1999) to the EMBL/GenBank/DDBJ databases.
- RC TISSUE=PLENTA;
- MELINE=2228825; PubMed=12477932; DOI=10.1073/pnas.242603899;
- RA Strauberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
- RA Klausner R.D., Colling F.S., Wagner L., Shemesh C.M., Schuler G.D.,
- RA Altschul S.F., Zeeberg B., Buettow K.H., Schaefer C.F., Bhat N.K.,
- RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh P., Diatchenko L., Marusina K., Farmer A., Rubin G.M., Hong L., Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Schatz T.E., Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
- RA Raha S.S., Loquaiello N.A., Peters G.J., Abramson R.D., Mulahay S.J., Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Guarante P.H., Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W., Villaon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A., Fahey J., Helton E., Ketteman M., Madan A., Rodrigues S., Sanchez A., Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G., Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C., Rodriguez A.C., Gimwood J., Schmutz J., Myers R.M., Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smailus D.E., Schnerer A., Schein J.E., Jones J.M., Marra M.A., and mouse cDNA Sequences.";
- RA Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
- [18]
- RP SEQUENCE OF 10-486 FROM N.A.
- RC TISSUE=Skeletal muscle;
- MELINE=96327611; PubMed=6672133;
- RA D'Esposito M., Quadreri N.A., Ciccodicola A., Bruni P., Esposito T., D'Urso M., Brown S.D.M.;
- RT "Isolation, physical mapping, and Northern analysis of the X-linked human gene encoding methyl CpG-binding protein, MECP2."; Mamm. Genome 7:533-535(1996).
- [19]
- RN SEQUENCE OF 10-486 FROM N.A.
- RA Reichwald K., Bauer D., Brenner V., Drescher B., Coy J.F., Kioschis P., Korn B., Nyakatura G., Platzer M., Poustka A., Sandhoff N., Rosenthal A.; Submitted (DEC-1996) to the EMBL/GenBank/DDBJ databases.
- RL [10]
- RP REVIEW ON VARIANTS.
- RP PubMed=12872250; DOI=10.1002/humu.10243;
- RA Miltenberger-Miltenyi G., Laccone F.; "Mutations and polymorphisms in the human methyl CpG-binding protein MECP2"; Hum. Mutat. 22:107-115(2003).
- RN [11]
- RP VARIANTS RTT TRP-106; CYS-133; SER-155; MET-158 AND CYS-306, AND VARIANT LYS-397.
- RP PubMed=10577905;
- RA Wan M., Lee S.S.J., Zhang X., Houwing-Manville I., Song H.-R., Amir R.E., Budden S., Naidu S., Pereira J.L.P., Lo I.F.M., Zoghbi H.Y., Schanen N.C., Francke U.; "Rett syndrome and beyond: recurrent spontaneous and familial MECP2
- RT mutations at CpG hotspots.";
- RL Am. J. Hum. Genet. 65:1520-1529(1999).
- RN [12]
- RP VARIANTS RTT TRP-106; CYS-133; SER-155 AND MET-158.
- RA MEDLINE=99438392; PubMed=10508514; DOI=10.1038/13810;
- RA Coy J. F., Sedlacek Z., Brechner D., Delius H., Poustka A.;
- RT "A complex pattern of evolutionary conservation and alternative polyadenylation within the long 3'-untranslated region of the methyl-CpG-binding protein 2 gene (MECP2) suggests a regulatory role in gene expression."; Hum. Mol. Genet. 8:1253-1262(1999).
- RL [13]
- RP INVOLVEMENT IN X-LINKED MENTAL RETARDATION WITH PROGRESSIVE SPASTICITY.
- RA PubMed=10986043;
- RA Meloni I., Bruttini M., Longo I., Mari F., Rizzolio F., Denvirleidt K., Fryns J.-P., Toniolo D., Renieri A.;
- RT "A mutation in the Rett syndrome gene, MECP2, causes X-linked mental retardation and progressive spasticity in males.";
- RL Am. J. Hum. Genet. 67:982-985(2000).
- RN [14]
- RP VARIANTS RTT VAL-100; GLN-105; TRP-106; CYS-133; ARG-152; SER-155; MET-158; ARG-305; CYS-306 AND HIS-306, AND VARIANTS CYS-86; MET-203; PRO-287; ALA-391; LYS-397; ILE-412 AND THR-444.
- RA PubMed=11055880;
- RA Buyse I.M., Fang P., Hoon K.T., Amir R.E., Zoghbi H.Y., Roa B.B., "Diagnostic testing for Rett syndrome by DHPLC and direct sequencing analysis of the MECP2 gene: identification of several novel mutations and polymorphisms.";
- RT Am. J. Hum. Genet. 67:1428-1436(2000).
- RL [15]
- RP VARIANT MRX16 VAL-140, AND VARIANT MET-203.
- RA MEDLINE=20465115; PubMed=11097980; DOI=10.016/S0014-5793(00)01994-3;
- RA Orrico A., Lam C., Galli L., Dotti M.T., Havek G., Tong S.F., Poos P.M., Zappone M., Federico A., Sorrentino V., "MECP2 mutation in male patients with non-specific X-linked mental retardation.";
- RL FEBS Lett. 481:285-288(2000).
- RN [16]
- RP VARIANTS RTT LEU-101; HIS-101; THR-101; TRP-106; CYS-133; CYS-134; ARG-152; MET-158; ARG-225; LEU-302; CYS-306 AND HIS-306, AND VARIANTS LEU-229 AND THR-439.
- RA PubMed=10767337; DOI=10.1093/hmg/9.7.1119;
- RA Cheadle J.P., Gill H., Fleming N., Maynard J., Kerr A., Leonard H., Krawczak M., Cooper D.N., Lynch S., Thomas N., Hughes H., Rulten M., Ravine D., Sampson J.R., Clarke A.; "Long-read sequence analysis of the MECP2 gene in Rett Syndrome patients: correlation of disease severity with mutation type and location.";
- RL Hum. Mol. Genet. 9:1119-1129(2000).
- RN [17]
- RP VARIANTS RTT GLN-105; MET-158; ARG-302; CYS-306 AND ALA-322.
- RA PubMed=10814711; DOI=10.1093/hmg/9.9.1377;
- RA Bienvenu T., Carrie A., de Roux N., Vinet M.-C., Jonveaux P., RA Couvert P., Villard L., Arzimanoglou A., Beldjord C., Fontes M., Tardieu M., Chilly J.; "MECP2 mutations account for most cases of typical forms of Rett syndrome.";
- RT Hum. Mol. Genet. 9:1377-1384(2000).
- RL [18]
- RP VARIANTS RTT MET-158; HIS-302 AND CYS-306, AND VARIANTS VAL-201; ALA-222; ILE-231 AND SER-376.
- RA PubMed=10944854;
- RA Amaro K., Nomura Y., Segawa M., Yamakawa K.; "Molecular analysis of the MECP2 gene in Japanese patients with Rett syndrome.";
- RT J. Hum. Genet. 45:231-236(2000).
- RL [19]
- RP VARIANTS RTT TRP-106; PHE-124; CYS-133; CYS-134; ARG-152; MET-158 AND CYS-306.
- RA MEDLINE=20439334; PubMed=10991688;
- RA Obata K., Matsushita T., Yamashita Y., Fukuda T., Kuwajima K., Mori K., Kondo I.;

RT "Mutation analysis of the methyl-Cpg binding protein 2 gene (MECP2) in patients with Rett syndrome.";
 RT J. Med. Genet. 37:608-610(2000).
 RL [20]
 RN
 RP VARIANTS RTT ARG-101; TRP-106; MET-158 AND CYS-306, AND VARIANT LYS-397;
 RX MEDLINE=2043935; PubMed=10991689;
 RA Hampson K., Woods C.G., Latif F., Webb T.;
 RT "Mutations in the MECP2 gene in a cohort of girls with Rett syndrome.";
 RT J. Med. Genet. 37:610-612(2000).
 RL [21]
 RP VARIANT RTT HIS-133.
 RX Published=11706982; DOI=10.1002/ana.1272;
 RA Armstrong J., Poo P., Pineda M., Aibar E., Gean E., Catala V.,
 RA Monros E.;
 RT "Classic Rett syndrome in a boy as a result of somatic mosaicism for a MECP2 mutation";
 RL Ann. Neurol. 50:692-692(2001).
 RN [22]
 RP VARIANTS RTT TRP-106; CYS-134; ARG-152; MET-158; ALA-302; CYS-306 AND ALA-322, AND VARIANTS VAL-201 AND LYS-397;
 RX Published=11738893; DOI=10.1016/S0381-7605(01)00342-4;
 RA Giunti L., Pegatti S., Lazzari V., Guarducci S., Lapi E.,
 RA Covillelo S., Cecconi A., Ombroni L., Andreucci E., Sani I.,
 Query Match 6.8%; Score 208.5; DB 1; Length 486;
 Best Local Similarity 23.9%; Pred. No. 0.00013; Matches 104; Mismatches 53; Indels 129; Gaps 17;
 . Db
 Qy 37 EDVAMELERYGEDEBQMMIKRSBCNPLQIPEPIASQFG-----ATACT 80
 :||: ;:::||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:
 22 KDKPKFKVKVKDKKKEKGKHEPVQPSAHHSAEAGAKETSEGSGSAPPVEASP 81
 . Db
 Qy 81 ECRKSV-----PCGWERVKQRLFGKTAGRFDVYFISPGQKFRSKSSLANY 127
 :||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:
 . Db
 Qy 82 KQRRIIIRDRGGPMYDPTLPGEWTRKLRQKGSRAGSKYDQVYLINPQKAFRSKVELAY 141
 :||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:
 . Db
 Qy 128 LHRNGETTSKPKEDPDFTVLKGKIGSKRYKOCMSAALTLHQNQSNNSNNMLRTSKCKD 187
 :||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:
 . Db
 Qy 142 FEKVGDTSLPNDPDFTV-TGRGSPSR-----REOKPPKK- 175
 :||:||:||:||:||:||:||:||:||:||:||:||:||:||:
 . Db
 Qy 188 VFMPPSSSLQESQLI-----SNFTSTHLLIKEDEGVDDVNFRKVPRKPKRVTILKGIPK 244
 :||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:
 . Db
 Qy 176 ---PKSPKAGFTGRGRGRPKGSQTTRPKATAGQEVQK-RVLRKSPK---LLVKMPF- 226
 :||:||:||:||:||:||:||:||:||:||:||:||:
 . Db
 Qy 245 KTKKGCRKSCSGFVQSDS-----KRESVNCNADAESEPAQKSQLDRTVCISDAGACGETL 300
 :||:||:||:||:||:||:||:||:||:||:||:||:||:
 . Db
 Qy 301 SVTSEBNLSVKKERNLSSGSNFCBOKTGGIINKPCSAKOSERNEKYEDTFLESEEIGT 360
 :||:||:||:||:||:||:||:||:||:||:||:
 . Db
 Qy 277 AAAABAKKAVKESSIR-----SVQETVLPIKK-----RKRET-----V 312
 :||:||:||:||:||:||:||:||:||:||:
 . Db
 Qy 361 KVEVVERKEHLHTDL-KRGSEMNONCSPTRKDGTGKTFQEDTPRQIERKTSLYP 418
 :||:||:||:||:||:||:||:||:||:||:
 . Db
 Qy 313 SIEVKEVKPKLVSTIGEKSGKGLKTCSPGRK-----SKESSPKGR----- 354
 :||:||:||:||:||:||:||:
 . Db
 Qy 419 SSKYNEALSPPRK 433
 :||:||:||:||:
 . Db
 Qy 355 ----SSSASSPPKE 365
 :||:||:||:||:
 RESULT 10
 ID 06QH19
 ID 06QH19 PRELIMINARY; PRT; 498 AA.
 AC 06QH19;
 DT 05-JUL-2004 (TREMBrel. 27, Created)
 DT 05-JUL-2004 (TREMBrel. 27, Last sequence update)
 DT 05-JUL-2004 (TREMBrel. 27, last annotation update)
 DE Methyl Cpg binding Protein 2 isoform B.
 Name=MECP2;
 Name=MECP2;
 Hom sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX PubMed=15034579; DOI=10.1038/ng1327;
 RA Matzakarian G.N., Lohi H., Munteanu I., Alfred S.E., Yanada T.,
 RA Macleod P.J., Jones J.R., Scherer S.W., Schanen N.C., Friez M.J.,
 RA Vincent J.B., Minassian B.A.;
 RT "A previously unidentified MECP2 open reading frame defines a new protein isoform relevant to Rett syndrome.";
 RT Nat. Genet. 36:339-341(2004);
 DR EMBL; AY541280; AAC55455.1; -;
 DR GO; GO:0005334; C:nucleus; IEA.
 DR GO; GO:003677; F:DNA binding; IEA.
 DR GO; GO:006555; P:regulation of transcription, DNA-dependent; IEA.
 DR InterPro; IPR000337; A+T hook.
 DR InterPro; IPR001739; Methy-Cpg bind.
 DR Pfam; PF02178; AT hook; 1.
 DR SMART; SP01428; MBD; 1.
 DR Pfam; PF01428; MBD; 1.
 SQ SEQUENCE 498 AA; 53323 MW; 443ECB3D5EA4DABB CRC64;

Query Match 6.8%; Score 208.5; DB 2; Length 498;
 Best Local Similarity 23.9%; Pred. No. 0.00014; Matches 104; Mismatches 149; Indels 129; Gaps 17;
 . Db
 Qy 37 EDVAMELERYGEDEBQMMIKRSBCNPLQIPEPIASQFG-----ATACT 80
 :||: ;:::||:||:||:||:||:||:||:||:||:||:||:||:||:
 . Db
 Qy 81 ECRKSV-----PCGWERVKQRLFGKTAGRFDVYFISPGQKFRSKSSLANY 127
 :||: ;:::||:||:||:||:||:||:||:||:||:||:
 . Db
 Qy 94 KQRRIIIRDRGGPMYDPTLPGEWTRKLRQKGSRAGSKYDQVYLINPQKAFRSKVELAY 153
 :||: ;:::||:||:||:||:||:||:||:||:||:
 . Db
 Qy 128 LHRNGETTSKPKEDPDFTVLKGKIGSKRYKOCMSAALTLHQNQSNNSNNMLRTSKCKD 187
 :||: ;:::||:||:||:||:||:||:||:||:
 . Db
 Qy 154 FEKVGDTSLPNDPDFTV-TGRGSPSR-----REOKPPKK- 187
 :||: ;:::||:||:||:||:||:||:||:
 . Db
 Qy 188 VFMPPSSSLQESQLI-----SNFTSTHLLIKEDEGVDDVNFRKVPRKPKRVTILKGIPK 244
 :||: ;:::||:||:||:||:||:||:||:
 . Db
 Qy 245 KTKKGCRKSCSGFVQSDS-----KRESVNCNADAESEPAQKSQLDRTVCISDAGACGETL 300
 :||: ;:::||:||:||:||:||:||:
 . Db
 Qy 239 QTSPGKKAEGGAGTTSTQWVNIKPGKRRKABDPQAIPIKR-----GRKPGSV 288
 :||: ;:::||:||:||:||:||:
 . Db
 Qy 301 SVTSEBNLSVKKERNLSSGSNFCBOKTGGIINKPCSAKOSERNEKYEDTFLESEEIGT 360
 :||: ;:::||:||:||:||:||:
 . Db
 Qy 289 AAAAAEAKKKAVKESSIR-----SVQETVLPIKK-----RKRET-----V 324
 :||: ;:::||:||:||:||:
 . Db
 Qy 361 KVEVVERKEHLHTDL-KRGSEMNONCSPTRKDGTGKTFQEDTPRQIERKTSLYP 418
 :||: ;:::||:||:||:||:
 . Db
 Qy 325 SIEVKEVKPKLVSTIGEKSGKGLKTCSPGRK-----SKESSPKGR----- 366
 :||: ;:::||:||:||:
 . Db
 Qy 419 SSKYNEALSPPRK 433
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 . Db
 Qy 367 ----SSSASSPPKE 377
 :||: ;:::||:||:

RESULT 11
 ID 072384 PRELIMINARY; PRT; 516 AA.
 AC 072384;
 DT 01-OCT-2003 (TREMBrel. 25, Created)
 DT 01-OCT-2003 (TREMBrel. 25, Last sequence update)
 DT 01-MAR-2004 (TREMBrel. 26, Last annotation update)
 DE Hypothetical protein DKFZp086A24160 (Fragment).
 Name=DKFZp086A24160;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 NCBI_TaxID=9606;

RN [1] SEQUENCE FROM N.A.
 RP TISSUE=Human colon endothelial primary cell culture;
 RC "Excessive hand-wringing in a MPPR-treated monkey.";
 RA Bloecker H., Brecher M., Mewes H.W., Weil B., Amid C., Osanger A.,
 RA submitted (JUN-2003) to the EMBL/GenBank/DDBJ databases.
 RLL EMBL; BX588060; CAD7991.1; -.
 DR HSSP; Q9UN29; IIG4.
 DR GO; GO:0005634; C:nucleus; IEA.
 DR GO; GO:0003677; F:DNA binding; IEA.
 DR GO; GO:0003555; P:regulation of transcription, DNA-dependent; IEA.
 DR SMART; SMO0391; MBD; 1.
 DR PFAM; PF01429; AT_hook; 1.
 DR SMART; SMO0391; MBD; 1.
 KW HYPOTHETICAL protein.
 FT NON_TER 1.
 SQ SEQUENCE 516 AA; 55204 MW; 27CD37B9164176B0 CRC64;
 Best Local Similarity 23.9%; Score 208.5; DB 2; Length 516;
 Matches 104; Conservative 53; Mismatches 149; Indels 129; Gaps 17;
 QY 37 EDVAMEELERGVEDEEQMMIKRSKSCCNLLQEFIASOFG-----ATAGT 80
 DB 52 KOKPLKKKKVKDKKEKEKGKHPVQPSAHSAAEPAKGAKBTSEGSGSAPAVPEASASP 111
 QY 81 ECRKSV-----PGGWERVQRQLFPTAQRFDVFISPOGLKRSKSLANY 127
 DB 112 KQRSRIRDRGPWMDDTLPLEGWTRKURKQKRSGRSAGKDYDVLINPQKARSKVLLAY 171
 QY 128 LHNGESELKPSDFDTFLSKRGIKSYKDCMAALTSHLQNSNNSNWNLRTSKKKD 187
 DB 172 FEKVGDSLDPNDPDFTV-TGKGSPLR-----REQKPKK- 205
 QY 188 VFMPPSSSELEQBSRGL--SNFTSHLLKEDEGVDDVNFRKVKKGKTYLKGPIK 244
 DB 205 ---PKSPKPAQGGRGRGRGPKGSQGTTPKAATSEGVOYK--RVLKSPGK--LLVKNPF- 256
 QY 245 KTKKGCKRCSGCVQSDS---KRESVNCNAKADAESEPAQKSQLDRTVCISDAGCETL 300
 DB 257 QTPGGKAEGGGATTSTQVMVTKRPGRKKAEDPQAIPKR-----GRKGSSVV 306
 QY 301 SVTSEESENLSVKKERSLSSGSNCSEOKTSGINGINKFCSAKDSBHNKEVDTPLESEBGT 360
 DB 307 AAAAAEAKKAVKESST-----SVOETVLPKK-----RKRET-----V 342
 QY 361 KVEVERKEHLHTDL--KRGSEMDNNCSPTRKDFTGKFIQFDTIIRTQERRKSLYF 418
 DB 343 STEVKVVKPLVSTLGKGLTKCKSPKR-----SKESSPKGR----- 384
 QY 419 SSKVNEALSPPRNK 433
 DB 385 ---SSSASSPPKE 395
 RESULT 12
 MEC2 MACPF STANDARD PRT; 486 AA.
 ID MEC2_MACPF
 AC 0951GB;
 DT 25-OCT-2004 (Rel. 45, Created)
 DT 25-OCT-2004 (Rel. 45, Last sequence update)
 DT 25-OCT-2004 (Rel. 45, Last annotation update)
 DE Match1-CPG-binding protein 2 (MECP-2 protein) (MECP2).
 GN Name=MECP2;
 OS Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Buteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;
 OC Cercopithecinae; Macaca.
 RN [1]

RN SEQUENCE FROM N.A.
 RP Muramatsu S.; "Excessive hand-wringing in a MPPR-treated monkey.";
 RC Submitted (AUG-2000) to the EMBL/GenBank/DDBJ databases.
 RA -!
 CC FUNCTION: Chromosomal protein that binds to methylated DNA. It can bind specifically to a single methyl CpG pair. It is not influenced by sequences flanking the methyl-CpGs. Mediates transcriptional repression through interaction with histone deacetylase and the corepressor SIN3A.
 CC SUBUNIT: Interacts with FNPBP3 (By similarity).
 CC SUBCELLULAR LOCATION: Nuclear. Colocalized with methyl-CpG in the genome (By similarity).
 CC SIMILARITY: Contains 2 A.T hook DNA-binding repeats.
 CC SIMILARITY: Contains 1 methyl-CpG-binding (MBD) domain.
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration between the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See <http://www.isb-sib.ch/announce> or send an email to license@isb-sib.ch).
 CC DR EMBL; AR25597; ARK97131.1; -.
 DR HSSP; P51608; IIG4.
 DR InterPro; IPR000637; AT_hook.
 DR InterPro; IPR001739; MeMyl-CpG_bind.
 DR Pfam; PF01429; MBD; 1.
 DR SMART; SMO0391; MBD; 2.
 DR PROSITE; PS50982; MBD; 1.
 KW DNA-binding; Nuclear protein; Repeat; Repressor; Transcription regulation.
 FT DOMAIN 90 162 MBD.
 FT DNA_BIND 185 197 A.T hook 1.
 FT DOMAIN 265 277 A.T hook 2.
 FT DOMAIN 366 372 His-rich.
 FT DOMAIN 376 405 Pro-rich.
 SQ SEQUENCE 486 AA; 52426 MW; 3471BB61d90d92A7D CRC64;
 Best Local Similarity 23.9%; Score 207.5; DB 1; Length 486;
 Matches 104; Conservative 53; Mismatches 149; Indels 129; Gaps 17;
 QY 37 EDVAMEELERGVEDEEQMMIKRSKSCCNLLQEFIASOFG-----ATAGT 80
 DB 22 KOKPLKKKKVKDKKEKEKGKHPVQPSAHSAAEPAKGAKBTSEGSGSAPAVPEASASP 81
 QY 81 ECRKSV-----PGGWERVQRQLFPTAQRFDVFISPOGLKRSKSLANY 127
 DB 82 KQRSRIRDRGPWMDDTLPLEGWTRKURKQKRSGRSAGKDYDVLINPQKARSKVLLAY 141
 QY 128 LHNGESELKPSDFDTFLSKRGIKSYKDCMAALTSHLQNSNNSNWNLRTSKKKD 187
 DB 142 FEKVGDSLDPNDPDFTV-TGKGSPLR-----REQKPKK- 175
 QY 188 VFMPPSSSELEQBSRGL--SNFTSHLLKEDEGVDDVNFRKVKKGKTYLKGPIK 244
 DB 176 ---PKSPKAPOTGRGRGRGPKGSQGTTPKAATSEGVOYK--RVLKSPGK--LLVKNPF- 226
 QY 245 KTKKGCKRCSGCVQSDS---KRESVNCNAKADAESEPAQKSQLDRTVCISDAGCETL 300
 DB 227 QTPGGKAEGGGATTSTQVMVTKRPGRKKAEDPQAIPIKR-----GRKGSSVV 276
 QY 301 SVTSEESENLSVKKERSLSSGSNCSEOKTSGINGINKFCSAKDSBHNKEVDTPLESEBGT 360
 DB 327 AAAAAEAKKAVKESST-----SVOETVLPKK-----RKRET-----V 312
 QY 361 KVEVERKEHLHTDL--KRGSEMDNNCSPTRKDFTGKFIQFDTIIRTQERRKSLYF 418
 DB 343 STEVKVVKPLVSTLGKGLTKCKSPKR-----SKESSPKGR----- 354
 QY 313 STEVKVVKPLVSTLGKGLTKCKSPKR-----SKESSPKGR----- 354

QY 419 SKYKNEALSPPRRK 433 DE Methyl-CpG-binding protein 2 (MeCP-2 protein) (MeCP2).
 ID : | | | | |::: GN Name=MeCP2;
 Db 355 ---SSSASSSPPKKE 365 OS Rattus norvegicus (Rat).
 OC Buxtorfa; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Murinae; Rattus.
 OX NCBI_TaxID=10116;

RESULT 13
 042403 PRELIMINARY; PRT; 344 AA.
 ID 042403; TISSUE=brain;
 DT 01-JAN-1998 (TREMBLrel. 05, Created)
 DT 01-JAN-1998 (TREMBLrel. 05, last sequence update)
 DT 01-OCT-2003 (TREMBLrel. 25, last annotation update)
 DB Attachment region binding protein (Fragment).
 GN Name=ARBP;
 OC Gallus gallus (Chicken).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Archosauria; Aves; Neognathae; Galiformes; Phasianidae; Phasianinae;
 OC Gallus.
 OX NCBI_TaxID=9031;
 RN [1] RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=7415642; PubMed=9271441;
 RA Weitzel J.M., Buhmester H., Straetling W.H.;
 RT "Chicken MAR-binding protein ARBP is homologous to rat methyl-CpG
 binding protein MeCP2";
 RL Mol. Cell. Biol. 17:5656-5666(1997).
 RN [2] RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=20188769; PubMed=10723722;
 RA Reichwald K., Theisen J., Wiatek T., Weitzel J., Straetling W.H.,
 RT "Comparative sequence analysis of the MeCP2-locus in human and mouse
 reveals new transcribed regions.";
 RL Mamm. Genome 11:182-190(2000).
 DR EMBL; Y1166; CAA74577.1; -.
 DR PDB; 1UB1; NMR; A=64-196.
 DR GO; GO:0005634; C:nucleus; IBA.
 DR GO; GO:000370; :transcription factor activity; IBA.
 DR InterPro; IPR000637; A+T hook.
 DR InterPro; IPR01739; Methyl-CpG-bind.
 DR InterPro; IPR00976; Wilms_tumour.
 DR Pfam; PF01429; MBD; 1.
 DR PRINTS; PR000929; ATHOOK.
 DR SMART; SM00391; MBD; 1.
 FT NON_TER 344 AA; 33640 MW; 53DD7BD9C9DF4FE3 CRC64;
 SQ SEQUENCE 344 AA;
 Query Match 6.6%; Score 202.5; DB 2; Length 344;
 Best Local Similarity 34.7%; Pred. No. 0.0002;
 Matches 59; Conservative 17; Mismatches 57; Indels 37; Gaps 5;
 QY 16 HAPTWTSERIVVDPDNPDLRKEDVAMLER----VGEDEEQMKTKRSSBCNPLIQLQEPITA 70 DE Query Match 6.6%; Score 201; DB 1; Length 492;
 DB 5 AAAAGGEGEELU---EEQADEGGVAGLKERPPKAKGKRKERRDEPAAEAEAPSGAEPAE 59 DE Best Local Similarity 22.9%; Pred. No. 0.00039; Mismatches 108; Indels 150; Gaps 18;
 QY 71 SAQFGATAGTEC-----RKS-----PCGWERVVKQRLFGKTG 104 DE Matches 108; Conservative 57; Mismatches 156; Indels 150; Gaps 18;
 DB 60 AGKADGSGGTAAPAVPEASAPKQKRSIRRDRGPMYDDTLPEGWTRKKQKRGSGAG 119 DE
 QY 34 LRKE-----DVAMELERVGEDBQMMIKRSSCNPILQEPITASAQFG----- 75 DE
 DB 8 LRKEKSEBDOLQGLKEPKLKPKVKKDKKEDBKGKHEPLOPSAHSABPAAGKAESSES 67 DE
 QY 76 -----ATAGTECRCSV-----PCGWERVVKQRLFGKTGAGRDWVFFSP 113 DE
 DB 68 SGSAAPAVPEASAPKQKRSIRRDRGPMYDDTLPEGWTRKKQKRGSGAGKVLYINP 127 DE
 QY 114 QGLKFRSKSSLANTLHKNGETSLKPKEDDPDTWLSKRGKSRVYKDCSMALTSILQNSNN 173 DE
 DB 128 QGKAFRSKVELIAYFEKQDLSLPNDPDDFTV-TGRGSPSR----- 167 DE
 QY 174 SNMLRLRTSKCKDVFMPSPSSBLOBSRGLSNFTSTLILKEDEGVDDNFRKV-RKPK 232 DE
 DB 168 -----REQKPK-----PKSKAPGTRGRGRGPKGSGTGRPKAASBQVQVKVLESP 217 DE

RESULT 14
 MEC2_RAT STANDARD; PRT; 492 AA.
 ID MEC2_RAT
 AC Q00566;
 DT 01-APR-1993 (Rel. 25, Created)
 DT 01-APR-1993 (Rel. 25, last sequence update)
 DT 25-DEC-2004 (Rel. 45, Last annotation update)

RESULT 15

MBC2_MOUSE STANDARD; PRT; 484 AA.

ID MBC2_MOUSE 99Z26; AC DT 16-OCT-2001 (Rel. 40, Created) 16-OCT-2001 (Rel. 40, Last sequence update)

DT 25-OCT-2004 (Rel. 45, Last annotation update)

DE Methyl-CpG-binding protein 2 (MeCP2 protein) (MeCP2).

GN Mus musculus (Mouse)

OS Mammalia; Eutheria; Rodentia; Chordata; Craniata; Vertebrata; Euteleostomi; Buka-Tyota; Metazoa; Sciuromorpha; Muridae; Murinae; Mus.

OC NCBI_TaxID=10090; OX RN [1]

SEQUENCE FROM N.A.

RC STRAIN=C57BL/6;

RX MEDLINE=9844942; PubMed=9774669; Hendrich B.; Bird A.; "Identification of a family of mammalian methyl-CpG binding proteins"; Mol. Cell. Biol. 18:6538-6547(1998).

RX MEDLINE=99399240; PubMed=10369871; DOI=10.1093/hmg/8.7.1253; Coy J. P.; Sedlacek Z.; Baechner D.; Delius H.; Pousetka A.; "A complex pattern of evolutionary conservation and alternative polyadenylation within the long 3'-untranslated region of the methyl-CpG-binding protein 2 gene (MeCP2) suggests a regulatory role in gene expression.;" Hum. Mol. Genet. 8:1253-1262(1999).

RN [3]

SEQUENCE FROM N.A.

RP REICHWALD K.; THIESSEN J.; WIEHE T.; KIOSCHIS P.; STRAETLING W.H.; ROSENTHAL A.; PLATZER M.; "Comparative analysis of the methyl CpG binding protein 2 locus in man and mouse reveals new untranslated sequences;" RIL Submitted (JUN-1999) to the EMBL/Genbank/DDBJ databases.

RN [4]

SEQUENCE FROM N.A.

RP STRAIN=FVB/N; TISSUE=Mammary gland; MEDLINE=22388257; PubMed=2477932; DOI=10.1073/pnas.242603899; RAX

RA STRAUBER R.L.; FEINGOLD E.A.; GROUVE L.H.; DERGE J.G.; KLAUBNER R.D.; COLLINS F.S.; WAGNER L.; SHENMER C.M.; SCHULER G.D.; ALCHUL S.R.; ZEEBORG B.; BUTSTOW K.H.; SCHAEFER C.F.; BHAT N.K.; HOPKINS R.F.; JORDAN H.; MOORE T.; MAX S.I.; WANG J.; HSIEH F.; BLATCHENKO L.; MARUSINA A.A.; FARMER M.; RUBIN G.M.; HONG L.; STAPLETON M.; SOARES M.B.; BONALDO M.F.; CASAVANT T.L.; SCHBEETZ T.E.; BROWN M.J.; USDIN T.B.; TOSHIBUKI S.; CARNICI P.; PRANGE C.; RHA S.S.; LOQUELLANO N.A.; PETERS G.J.; ABRAMSON R.D.; MULILARY S.J.; BOSEK S.A.; McEWAN P.J.; MCKERNAN K.J.; MALEK J.A.; GUARANTE P.H.; RICHARDS S.; WORLEY K.C.; HAILE S.; GARCIA A.M.; GAY L.J.; HULYK S.W.; VILLALON K.; MUZNY D.M.; SODERGREN E.J.; LU X.; GIBBS R.A.; PHEY J.; HEITON E.; KETTEMAN M.; MADAN A.; RODRIGUES S.; SANCHEZ A.; WHITING M.; MADAN A.; YOUNG A.C.; SHEVchenko Y.; BOUFFARD G.G.; BLAKESLEY R.W.; TOUCHMAN J.W.; GREEN B.D.; DICKSON M.C., RA

RA Rodriguez A.C.; GRIMWOOD J.; SCHMUTZ J.; MYERS R.M.; SMITHUS D.E.; RA BUTTERFIELD Y.S.N.; KRZYWIŃSKI M.I.; SKALSKA U.; SMAILUS D.E.; RA SCHNEIDER A.; SCHEIN J.E.; JONES S.J.M.; MAZRA M.A.; RA "Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences;" Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002). RA RN [5]

RRP INTERACTION WITH FNBP3. MEDLINE=97315177; PubMed=9171351; DOI=10.1093/embj/16.9.2376; RR BEDFORD M.T.; CHAN D.C.; LEDER P.; "FNBP WW domains and the Ab1 SH3 domain bind to a specific class of proline-rich ligands;" EMBO J. 16:2376-2383 (1997).

CC -1- FUNCTION: Chromosomal protein that binds to methylated DNA. It can bind specifically to a single methyl-CpG pair. It is not influenced by sequences flanking the methyl-CpGs. Mediates transcriptional repression through interaction with histone deacetylase and the corepressor SIN3A (By similarity).

CC -1- SUBUNIT: Interacts with FNBP3.

CC -1- SUBCELLULAR LOCATION: Nuclear. Colocalized with methyl-CpG in the genome.

CC -1- SIMILARITY: Contains 2 A-T hook DNA-binding repeats.

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CC DR EMBL; AF07251; AAC68880; 1; -. DR EMBL; AF15911; AAC77153; 1; -. DR EMBL; BC02153; AAH77153; 1; -. DR HSSP; P51608; 1QK9.

DR MGD; MGI:99918; MeCP2.

DR GO; GO:0005634; C:nucleus; IDA.

DR InterPro; IPR000637; A+T hook.

DR InterPro; IPR001739; MeChyl-CpG_bind.

DR PF01429; MBD; 1.

DR SMART; SM00384; AT hook; 2.

DR PROSITE; PS50982; MBD; 1.

KW DNA-binding; nuclear protein; repeat; Repressor; transcription regulation.

FT DOMAIN 90 162 MBD.

FT DOMAIN 185 197 A-T hook 1 (By similarity).

FT DOMAIN 265 277 A-T hook 2 (By similarity).

FT DOMAIN 366 372 His-rich.

FT DOMAIN 379 403 Pro-rich.

SO SEQUENCE 484 AA; 52307 MW; 62FU228F0118A49F CRC64;

Query Match 6.5%; Score 198.5; DB 1; Length 484; Best Local Similarity 24.1%; Pred. No. 0.00054; Matches 91; Conservative 50; Mismatches 154; Indels 83; Gaps 11;

QY 76 ATAGTCRKSV-----PGCWERVKQRKGKTAGRFDWYFISPOGLKRSKS 122

Db 77 ASASPKQRKSRIRDGRGMYDDETLPPGWTRKKQRKRSAGKYDVYLNGOKAFRSKV 136

QY 123 SLANYLKHGETSLKEPFDVTLSRKGKIKYKDCCMAALTSHLQNQSNNSNWLRTRS 182

Db 137 ELJAYFKVQGLSDPDPDFTY-TGCGSPREQ----KPKKPKSPKAPGTRGRG 189

QY 183 KCKKDVMPSSSSELQBSRGLSNFTSHLILKEDEGVDDYNFRKVPKPK-----VT 236

Db 190 R-----PKGSITGRPKAASEGIVQVKRLKSKVLUVKMPFCASPGKGEGGATT 241

QY 237 IKGKIPKTKKGCRKCSGKVQDSKRE----SYCNKADAESPVAKS---QLDRT 287

Db 242 SAQVNVIK--RPGRKRAEDPOAIPKKRGKPGSVAAAEAKKAVKESSIRSVHET 299
Qy 288 VCISDAGACSETLVTSEE-----NSUVKKERSLSGGSNFCSBQKTSGLINKFCSAK 340
Db 300 VLPPIKRKRTRTIVSIEVKVWKLVLVSTLGKSGKLKTCSPGRKSKESSPKERSSAS 359
Qy 341 DSEHNNEKYEDTFLESBEIGTKVEVVERKERHLHTDILK-----RGSEMDDNNCSP 388
Db 360 SPPKUEHHH-----RHHHSESTIKAPMPLPSPPPPEPESSEDPISP 400
Qy 389 TRKDFGTGEKLFQEDTIPR 406
Db 401 PEPQDLSSSSICKEEMPR 418

Search completed: August 22, 2005, 10:09:03
Job time : 182 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 22, 2005, 10:02:08 ; Search time 42 seconds
(without alignments)
1328.708 Million cell updates/sec

Title: US-10-629-951-2
Perfect score: 3055
Sequence: 1 MGTIGLSSLSLGDRGANPTV..... HKLNKYHDWLWENHEKLSLS 580
Scoring table: BLOSUM62
Gappen 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR 79,*

1: pir1;*
2: pir2;*
3: pir3;*
4: pir4;*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	209	6.8	476	2 S57963	methyl Cpg binding
2	6.6	492	2 A41907	methyl-CpG-binding	
3	4.7	1301	2 S51323	SAC3 protein - yeast	
4	4.7	348	1 G02334	breast cancer tumor	
5	4.6	2	27833	rhoptry-associated	
6	4.5	500	2 S55785	nucleolar protein	
7	4.5	560	2 S53382	protein YKR029c homolog	
8	4.5	598	2 B40713	cyclin I - human	
9	4.5	1359	2 T34036	hypothetical protein	
10	3.5	4.4	822	T46262	probable ABC transporter
11	3.2	4.3	286	A77682	probable A/G-specific
12	3.2	4.3	1819	2 A71928	cag island protein
13	3.0	4.3	997	2 T43523	cut117 protein - filaggrin
14	3.0	1927	2 G65829	pathogenicity	
15	2.8	4.2	946	2 A96748	hypothetical protein
16	2.8	4.2	853	T51505	hypothetical protein
17	2.6	4.1	561	2 H86442	unknown protein [1]
18	2.5	4.1	491	2 T50346	hypothetical protein
19	2.4	4.1	650	2 T33550	hypothetical protein
20	2.4	4.1	1702	2 T14050	protein kinase
21	2.3	4.0	697	2 C97120	topoisomerase I [1]
22	2.2	4.0	669	S55024	ribulin, skeletal
23	2.2	4.0	651	2 C86333	hypothetical protein
24	2.2	4.0	3924	2 S37431	ankyrin 2, neurona
25	2.0	3.9	911	2 S54491	hypothetical protein
26	2.0	3.9	1147	2 JN0599	DNA-binding protein
27	2.0	3.9	1233	2 S56271	hypothetical protein
28	2.0	3.9	1131	A49393	activator 1 large
29	2.0	3.9	1440	2 T33813	hypothetical protein

ALIGNMENTS

Qy	37 EDVAMIELSERVGEDESOMWMKRSSECNPILQBLPIASAQFG-----ATAGTE 81	Db	13 KDKPLKEKKVKQKDKKEKEKGKHPVQPSAHSAAEPAGRAETSEBGSGARLCEASASPK 72
Qy	82 CRKSV-----PCGWERVVKQRLFEGTKAGRDFYFISQGLKRSKSSLANVL 128	Db	73 QRRSIRDGRPMWDDPPLTPECLPSEGWTKLKQKRSRGSAGKSYQDYLINFGKATRKSKVLLAYF 132
Qy	129 HKNGERTSLKPSDDFTLUSKQGKIKSYKDCSMAALTSHLQNQSNNNNLURTRSKCKDV 188	Db	133 EKVGDISLDPNDFDFTV-TGRGSPSR-----REQPKPK-- 165
Qy	189 FNPSSSSELOBSRGI-----SNFTSTILLKEDEGVDDVNFRKVRKPKGVMTLGKIPKK 245	Db	166 ---PKSKPAGTAGTGRGRGRPKGSQGTTPKRAATSEGIVQVK--RVLKSPGK--LLVKMPF-Q 217
Qy	246 TKKGCRKSCSGFVQVSDD---KRESVCKNADAESEAQAQSQQLDRVCISDAGAGETLS 301	Db	218 TSPGGKAGEGATTSTQWVTKRPSRKAKADPAPKPK-----GRKPSVVA 267
Qy	302 VTSBENSILVKKERSLSSGSNPCEQRTGSGINKFKCSAKOSBNEKEVDPLESEBIGTK 361	Db	304 IEVKEVVKPLLVSTLGKSGKGKLTCKSPGRK-----SKESSPKGR----- 344
Qy	420 SKYNKEALSPPRK 433	Db	345 ---SSSSSPRKE 355

RESULT 2

A;Accession: A41907
methyl-CpG-binding protein 2 - rat
N;Alternate names: chromosomal protein MeCP2
C;Accession: A41907; S4461
R;Lewis, J.D.; Meehan, R.R.; Henzel, W.J.; Maurer-Fogy, I.; Jeppesen, P.; Klein, F.; Birrell, G.; Purification, sequence, and cellular localization of a novel chromosomal protein A;Reference number: A41907; MUID:92298389; PMID:1606634
A;Accession: A41907
A;Status: preliminary
A;Molecule type: mRNA
A;Residues: 1-492 <LEW>
A;Cross-references: UNIPROT:Q00566; GB:N94064; NID:9205361; PIDN:AAA41584.1; PID:9205362
R;Nan, X.; Meehan, R.R.; Bird, A.
Nucleic Acids Res. 21, 4886-4892, 1993
A;Title: Dissection of the methyl-CpG binding domain from the chromosomal protein MeCP2.
A;Reference number: S41611; MUID:9423813; PMID:8177735
A;Contents: annotation; methyl CpG-binding domain
C;Keywords: chromosomal protein; DNA binding
F:75-162/Domain: methyl-CpG-binding #status experimental <MCG>

Query Match 6.6%; Score 201; DB 2; Length 492;
Best Local Similarity 22.9%; Pred. No. 2.2e-05; Mismatches 156; Indels 150; Gaps 18;
Matches 108; Conservative 57; MisMatched 156; Indels 150; Gaps 18;

Qy 34 IRKE-----DVAMELERVEDEEONMIKRSSECNFILLOPIATSAQFG----- 75
Db 8 LERKEKSEDQDQLGLKEKPLKEKKVKKDKKEKGKHPELQPSAHHSAPFAAGAKTSES 67

Qy 76 -----ATAGTECRKV-----PGWERTVKQRLFGKTAGRFVYTFSP 113
Db 68 SGASAPAVPEASPKORRSLDRGPMWYDPTLPKGWTTRKQKRSIGRSAGKYDYLINP 127

Qy 114 OGLKERSKSSAIALNYLHKNGETSLKQEDFDITVLSLRGKIKRSYKDCSMAALTSHLQNQNN 173
Db 128 OGKAFRSKVELLAVYFVKGTSLSLDPMDPFTV-TGRGSFSR----- 167

Qy 174 SWNNLNRTRSKCKDVFMPRSSSSELQBSRGSNSNFTSHLLIKEDEGVDDVNFRKVY-RKPK 232
Db 168 -----REQKPKK-----PKSKPAPGTGRGRPKSGTGPKRPAKALASEGVQVQRVLEKSP 217

Qy 233 GKVTILKGIPKTKKGCRKCSGFGVQSDS-----KRESYCNKADAESEVAQSQLDRT 287
Db 218 GKR-LIVKMPFQASPG---KOGGGATTTSQAWMVTGPKRKAQADPOQIIPKGR----- 268

Qy 288 VCISDAGACGETLSTVSEENSLVKKERSLSSGSNFCSEOKTISLINKFCSSAKOSENEK 347
Db 269 -----GRKPSSWAAAABAKKAVKESIR-----SYQETVLLIKK-----RK 307

Qy 348 YEDTTILESEEITGKTYEVVERKEHLHTDL--KRGSEMDNNCSPTKRDFTGCKIFQEDTIP 405
Db 308 TRET-----VSIEVKVVKPLLUSTLGKSGKLCKSPGRK-----SKESSP 351

Qy 406 RTQIQRKTSLYFSSKYNKELSPRR-----KAFKKWTPPRP 444
Db 352 KGR-----SSSSSPKKEHHHHHAESPCKAPMPLPPPPP 388

RESULT 3

S51323 SAC3 protein - yeast (Saccharomyces cerevisiae)
N;Alternate names: protein YDR358.13; protein YDR159w
C;Species: Saccharomyces cerevisiae
C;Date: 23-Feb-1995 #sequence_revision 12-May-1995 #text_change 09-Jul-2004
C;Accession: S51323; S55983; S71744
R;Bauer, A.; Koelling, R.
submitted to the EMBL Data Library, January 1995
A;Description: The SAC3 gene codes for a nuclear protein required for normal mitosis.
A;Reference number: S51323

A;Accession: S51323
A;Molecule type: DNA
A;Residues: 1-1301 <BAU>
A;Cross-references: UNIPROT:P46674; EMBL:247805; NID:9634085; PID:9634086
A;Cross-references: UNIPROT:P46674; EMBL:247805; NID:9634085; PID:9634086
R;Murphy, L.; Richards, C.; Harris, D.
submitted to the EMBL Data Library, July 1995
A;Reference number: S51323
A;Accession: S57983
A;Cross-references: EMBL:Z50046; NID:9899393; PID:9899406; MIPS:YDR159w
A;Cross-references: EMBL:Z50046; NID:9899393; PID:9899406; MIPS:YDR159w
R;Bauer, A.; Koelling, R.
Yeast 12, 965-975, 1995
A;Title: Characterization of the SAC3 gene of *Saccharomyces cerevisiae*.
A;Reference number: S71744; MUID:97027306; PMID:8873450
A;Cross-references: SGD:S71744
A;Molecule type: DNA
A;Accession: S71744
A;Cross-references: SGD:S71744
A;Genes: SGD:SAC3; LEP1
A;Cross-references: SGD:S0002566; MIPS:YDR159w
A;Map Position: 4R
C;Function:
A;Description: potential regulator of leucine permease gene expression
C;Keywords: nucleus; transmembrane #status predicted <TM>
F:999-1015/Domain: transmembrane #status predicted <TM>

Query Match 4.7%; Score 115; DB 2; Length 1301;
Best Local Similarity 21.9%; Pred. No. 0.32; Mismatches 185; Indels 168; Gaps 25;
Matches 116; Conservative 61; MisMatched 185; Indels 168; Gaps 25;

Qy 123 SLANVTHKNGETSLKQEDFDITVLSLRGKIKRSYKDC-----KDFVNPSSSBLOSRLSNFTSHL 212
Db 510 TLQMSHAKLSETOBQLKKY-LTICERQRQKTRYKGLINGEDNLASSYYVKDPKKDRIPS 568

Qy 160 MAALTSHLQNOSINNNWNLRTSKCK-----KDFVNPSSSBLOSRLSNFTSHL 212
Db 569 IADQSFMLMENFQNTYNEKINONNSVYKPOINTSPKRVATRPNHPPFSQSKOLPQISQHT 628

Qy 213 LKKEDEGVDDVNFRKVPKRKGKVYVTLKQGKPIKKKGCRKSCSGFVQDSKRSKRSVCNRA 272
Db 629 L-----STNPILITPVQHGDSLSEQQQIKTVTDG----GSPFVFDQSAQNSTVEASK 676

Qy 273 AESEPVQASQLDRTVCTS DAGAGGETSVTSSENSLVKKKENSUSSSNFCSEOKTSGI 332
Db 677 A-----HMISTVSGANGDEKUS---SBOEMERKEI-----EEKTC----- 712

Qy 333 INKFCSAKUDSEHNENKEYDTFLESBEIGKV--EVVERKEHLHTDLKGSEMNDNCSPTR 390
Db 713 -----QLKCKQEN--ADQKVITEQIANDLVKUVNNSV--ISIVKEFSBAN----YR 757

Qy 391 KDFVQEKIFQEDTIPR---TQERRKTSLYFSSKYNKELSPPRKAFKKW----- 438
Db 758 KDFI-----DTWTRELYDAFLHERLYLIMDSRAEJKRNSTKKFFEWQASQK 810

Qy 439 -----TPPRSPNLUQV-----TLEFDPWKLIAITFLNRSGKMAI--PY-- 477
Db 811 KNRLIBEKKEEIKVSLQVGPFKKTCFLFRPYKVNNSFMLSSDKNLFPSPLND 870

Qy 478 -----LWKFLF-----KVPSEAERTADWRDVSLIKPLGILDR 512
Db 871 EFNKFATHLTFLKISKWLRPLMQSIYDNLTKKPF-----SNSLTANLF--IY 916

Qy 513 AKTIVKFSDSYLTLKQWCKVPIELIGIGKQND--SYRIFCVNEWKQVHED 560
Db 917 AKDWTSLSNRWILSKFNLO-TAQDSKKSNNISSRICCIDD--EYESD 963

RESULT 4

G0234 breast cancer tumor suppressor BRCA2 - human

N;Alternate names: breast cancer susceptibility protein BRCA2
 C;Species: Homo sapiens (man) sequence_revision 06-Jun-1997 #text_change 09-Jul-2004
 C;Date: 21-Dec-1996
 C;Accession: G02334; S88501
 R;Tavtigian, S.V.; Rommens, J.M.; Couch, F.J.; Neuhäusen, S.; Bell, R.; Berry, S.; Bogde
 er, M.; Snyder, S.; Stringfellow, M.; Stroup, C.; Swedlund, B.; Teng, D.; Thomas, A.; Ti
 submitted to the EMBL Data Library, December 1995
 A;Reference number: H01078
 A;Status: translated from GB/EMBL/DDJB
 A;Molecule type: mRNA
 A;Residues: 1-3418 <TA>
 A;Cross-references: UNIPROT:P51587; EMBL:U43746; NID:91161183; PIDN:AB07223.1; PID:9116
 R;Woosier, R.; Biagini, G.; Lancaster, J.; Swift, S.; Seal, S.; Mangon, J.; Collins, N.;
 ith, A.; Connor, P.; Arason, A.; Guðmundsson, J.; Ficenec, D.; Kelsell, D.; Ford, D.;
 ; Narod, S.; Deno, G.; Egilsson, V.; Barkadottir, R.B.; Baston, D.F.; Bentley, D.R.
 Nature 378, 789-792, 1995
 A;Authors: Futreal, P.A.; Ashworth, A.; Stratton, M.R.
 A;Title: Identification of the breast cancer susceptibility gene BRCA2.
 A;Reference number: S68501; NID:96112016; PMID:8524414
 A;Molecule type: mRNA
 A;Residues: 282-371, 'N', 373-598, 'S', 600-1108, 'EQ', 1111-1119, 'D', 1121-2321, 'V', 2323-2386,
 C;Genetics:
 A;Cross-references: GDB:387848; OMIM:600185
 A;Map position: 13G12.3-13q12.3
 C;Superfamily: DNA recombination repair protein, BRCA2 type
 C;Keywords: polymorphism; tumor suppressor

Query Match 4 7%; Score 145; DB 1; Length 3418;
 Best Local Similarity 20.9%; Pred. No. 1.2; Matches 96; Conservative 67; Mismatches 158; Indels 138; Gaps 23;
 Qy 36 KEDVWAMELERVGEDBQMMIKRSKSCNPLQOPIQSQAQFGMTAGTTCRKSVPFGWRRVK 95
 Db 972 KSDPSLNIDKPEKONDYMKWAG---LIGPISNSPGCSFRTRASNEKEIQLSEINIKK 1026
 Qy 96 QRLFEKTAGRFDPDVYFISPOGLKFRSKSLANYLHKINGETSL-KPBDPDFTYLSKRIGIKSR 154
 Db 1027 SKMFFK----DIEBOYPTSL---ACVEIVNTLADNOKKLSKPOSIN-TVSAHQSSV 1077
 Qy 155 YKDCCSMAALTSHL--ONQSNNSNNWILTRSKCKDQVMPMPSSSEIOPERSGLSNPSTHIL 212
 Db 1078 VSDCCKNSHTTPQMLSKQDFNHNH-----TBSQAKETE-----LSTIL 1118
 Qy 213 LIKPEREGVDDVNFRKYR-----PKGKVILK----- 239
 Db 1119 ----PESGSQFETPQRKPSYTLQKGSTPEVENQNMILTKTSBECRADLHVIMMAPSIG 1174
 Qy 240 -----GIPKK-----TKGGRKSCSGFVQSDSKRSCRVCKNDAESEPVAKSQI 284
 Db 1175 QVDSSKQFEGTWEIKRKFKAGLKLNDCNKSASGYLTDR-----NEVGPRGPFYSAHGTLK 1227
 Qy 285 D-RTWVTSIDAGACGETLSVTBENSILVKKERSLSSGSNFCSBQKTSGLINKFCASKDSE 343
 Db 1228 NVSTEAHQKAVKLUFSPIENISBTS---AEVPHPSLSSKCD---SVVSNP-----KLEN 1277
 Qy 344 HNEKVKHDTFLISBEE-----IGTKVEVERKEHLTDILKRGSEMDDN-CSP 388
 Db 1278 HRDK---TVSKKNCQCOLLQNNIEMTGIFVTEI-----TENYGRANTEENDKNTAA 1327
 Qy 389 TRK---DPFGKFKQBDTIPRTOERRKTSYPPSKY 423
 Db 1328 SRNHSNLEFDGSDSKNDTV---CIIKDETLIPTDORN 1363

RESULT 5
 S27033
 rhoptry-associated protein 1 precursor - malaria parasite (Plasmodium falciparum)
 N;Alternate names: protective antigen
 C;Species: Plasmodium falciparum
 C;Date: 17-Apr-1993 #sequence_revision 17-Apr-1993 #text_change 09-Jul-2004

C;Accession: A45514; S27833
R;Ridley, R. G.; Takacs, B.; Lahn, H. W.; Delves, C. J.; Goman, M.; Certa, U.; Matile, H.; Mol; Bloch, Mol. Biochem. Parasitol.; 41, 125-134, 1990
A;Title: Characterisation and sequence of a protective rhoptry antigen from Plasmodium A;Reference number: A45514; MUND:90348711; PMID:2200961
A;Accession: A45514
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-782 <R12>
A;Cross-references: UNIPROT:Q26007; GB:M32853; NID:9160656; PID:9160657
C;Super-organisms: Plasmodium falciparum rhoptry-associated protein 1

A;Accession: S6805; A;Status: nucleic acid sequence not shown; not compared with conceptual translation
 A;Molecule type: mRNA
 A;Residues: 374-407 <VAN>
 R;Brown, D.; Churcher, C.M.; Barrell, B.G.; Rajandream, M.A.; Wood, V.
 submitted to the EMBL Data Library, September 1997
 A;Reference number: Z21733
 A;Accession: T37634
 A;Status: preliminary; translated from GB/EMBL/DDJB
 A;Molecule type: DNA
 A;Residues: 1-338; 'S', 340-500 <BRO>
 A;Cross-references: EMBL:Z99091; PIDN: CAB1172-1; GSDB: GN00066; SPDB: SPAC13F5.09
 C;Genetics:
 A;Gene: gar2; SPDB: SPAC13F5.09
 A;Map position: 1
 C;Superfamily: ribonucleoprotein repeat homology <RRM1>
 F:264-331/Domain: ribonucleoprotein repeat homology <RRM2>
 Query Match 4.5%; Score 138.5; DB 2; Length 500;
 Best Local Similarity 23.7%; Pred. No. 0.23; Mismatches 147; Indels 41; Gaps 11;
 Matches 74; Conservative 50; Mismatches 147; Indels 41; Gaps 11;
 Qy 185 KKDVFMPSSSELEQESRLSNFTSHILKEPGDDUNFRKTKPKGKVTLKGIPK 244
 Db 19 KKGAIKEKSKKTKEAAK--IAKQSKTDVSPKSKKEAKRASSPE-PSK 68
 Qy 245 KTKKGCKRSCKSGFVQDSKRESVONKADEAEPVQKQSOLDRTVCIISAGACETLSVT 304
 Db 69 KSTVKQKSKK--KEESESSSESSSESSSESSSESSSESSSESSSESSSESSSESSSESS 122
 Qy 305 EENSLVK--KKERSLSSGSNFCSQTKSGIINFKFCASDKESENEKYEDTFLESEIGK 361
 Db 123 BEEVIVTKBKKSESSSESSSESSSESEEEBAV-KIEEKKESSSDSSSESSSESSSESS 181
 Qy 362 VEVVERKELIH-TDLIKGSEMNONCSPRKDFEGKIFOEDTPRTOLEKKTSLYSS 420
 Db 182 SESBEEEEEVEKTEKEKKGSSSESSSDHSSSDSSSE--SGDSDSSDSESESS---SE 234
 Qy 421 KYNKEALSPPRRAFKKWTPROPFNLVQETLHDWKLIATIFLNRTSGKMAIPVLMK 480
 Db 235 DEKKRKAASPASERPAKTTKPSODSN--ET-----CTWFVGULSWNTDQWLGQ 281
 Qy 481 FLERKVPSEAVR 492
 Db 282 EFFEYGTIVGAR 293

RESULT 7

553382
 protein YKR029c homolog YJH105w - Yeast (Saccharomyces cerevisiae)
 N;Alternate names: hypothetical protein J0819
 C;Species: Saccharomyces cerevisiae
 C;Date: 05-May-1995 #sequence revision 01-Sep-1995 #text_change 09-Jul-2004
 C;Accession: S53382; S56883; S57363
 R;Rasmussen, S.W.
 submitted to the EMBL Data Library, February 1995
 A;Description: A 37.5 kb region of yeast chromosome X includes the SME1, MEF2, GSH1
 A;Reference number: S53376
 A;Accession: S53382
 A;Molecule type: DNA
 A;Residues: 1-560 <RAS>
 A;Cross-references: UNIPROT:P42948; EMBL:X85021; NID:g728698; PID:g728705
 R;Rasmussen, S.W.
 submitted to the Protein Sequence Database, September 1995
 A;Reference number: S56876
 A;Accession: S56883
 A;Molecule type: DNA
 A;Residues: 1-560 <RAW>
 A;Cross-references: EMBL:Z49380; NID:g1008285; PID:g1008286; MIPS:YJL105w
 R;Rasmussen, S.W.
 yeast, II, 873-883, 1995

A;Title: A 3'-5' KD region of Yeast cdc200cogene A includes the Shizuka, Maruyama, Saito and Yodoi
A;Reference number: S557357; PMID:96090136; PMID:7483851
A;Accession: S557363
A;Sequence: nucleic acid sequence not shown; translation not shown
A;Molecule type: DNA
A;Residues: 1-560 <RAF>
A;Cross references: EMBL:X85021; NID:9728698; PIDN:CAA59389_1; PID:9728705
A;Note: the nucleotide sequence was submitted to the EMBL Data library, February 1995
C;Genetics:
A;Cross-references: SGD:S0003641
A;Map position: 10L

QY	174 SNHNLRTRSKCKD---VMPRSSSSESQES-----RGLSNFTSHLILKEDBEGVD	Db	1344 -----EVKSVKAVLDCCVSARBNRKERKECEKLIT----- 1373
QY	587 SEOSLQLLSESENDDKLPLIPLAIKKDKDNLVGSLEKGKSTSK--TKFDISIVD	Db	273 ABSEPVQK----SOLDRV--CI----SDAGACGETLSVSEENSUVKKERSLSS 319
QY	223 VNTRKVPRPKGKOTLKGKPIKTKKGKRSKSGFVQ-----SDSRKESV 267	Db	1374 ---PEARFLAKELQDKDKAICDKLNADPNRRAIMKCLDGSLBEKL----- 1419
QY	644 P---IEKPTTEIS---EVLPBEECRKAICDQSFRVRSTDGRVTKTRDVSSPVSDEKENV 697	Db	320 GSNPCSBQTKGTINPKCSAKUSENREKVTITLE-SEE-GTKEVERGEHLHDILR 378
QY	268 CNKDAES-----EVAQSKQLDRTVICSDA--GACGETLSVSEBNLSVK 312	Db	698 -NHREANSGHTVMVNHHSLDPQIVPMELEGSGSYLQKLPDRNVMNGSERKVTFQEDINSP 756
QY	313 KERLSSGSNCBQKTSGIINPKCSAKDSEHNEKYETFLFLESBEGTKEVERKHLH 372	Db	379 GSE-MDNCSPRKD 392
QY	757 KLOSKNOTVEANETSDKLOSKKEAHLENLEKIEEKLTEVDKVSLSDAFPDQEIKNS 816	Db	373 TDILKGSEBMNDNCSPTRKDFTRKIFQBDTIRTOERR---KTSLYFSKX-INTERL 427
Db	817 RTSYQNGTRSVSQNTPEKE---TKVDKIDUNVERKDKVETSPSPSCETSSAFAKTYAEKVT 872	Db	428 S---PPRKAKFKK--WTPRSPFLNUVOETLHPWKLLIATFLNRTSGKMAIPVNLKEL 482
QY	873 SINPSVVKPLDSEYYDHSISPDPLCQSSFLP-----QTPVSKHALPLV---- 919	Db	483 EKYPSEAVARTADWRR--DVSELK---PLGYLDRLAKTIVKESDEVILKOM-KYTEL 534
QY	920 -----EANAPPWERPIDSLLSPVNPVPENKLSKEL---DMTVEQMIKP---- 963	Db	920 -----EANAPPWERPIDSLLSPVNPVPENKLSKEL---DMTVEQMIKP---- 963
QY	535 HGIGKKGDSYRIFCVNEMKQVHEDHKLANKYHDWLWNHEK 576	Db	964 ---MYAK---CAKEFEAAEE--KL---EWLLEBGKR 989
QY	964 ---MYAK---CAKEFEAAEE--KL---EWLLEBGKR 989	Db	964 ---MYAK---CAKEFEAAEE--KL---EWLLEBGKR 989
RESULT 14			
GE4585	cag pathogenicity island protein cag7 - <i>Helicobacter pylori</i> (strain 26695)	Db	A9674B ----- 1
C.Species:	<i>Helicobacter pylori</i>	Db	A9674B ----- 1
C;Accession:	GE4585	Db	A9674B ----- 1
C;Accession:	90-Aug-1997 #sequence_revision 09-Aug-1997 #text_change 09-Jul-2004	Db	A9674B ----- 1
R.Theologis, A.; Becker, J.R.; Palm, C.J.; Federpiel, N.A.; Kaul, S.; White, O.; Chin, C.W.; Chung, M.K.; Conn, B.; Conway, A.R.; Creasy, T.H.; Dews, N.F.; Hughes, B.; Huizar, L.	Db	A9674B ----- 1	
Nature 408, 816-820, 2000	Db	A9674B ----- 1	
A;Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, S.X.; Liu, Z.A.; Luos, J.S.; Maiti, R.; Matz, M.; Rooney, T.; Rowley, D.; Sakano, H.	Db	A9674B ----- 1	
A;Authors: Salzberg, S.L.; Schwartz, J.R.; Shin, P.; Southwick, A.M.; Sun, H.; Turner, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.	Db	A9674B ----- 1	
A;Title: Sequence and analysis of chromosome 1 of the plant <i>Arabidopsis</i> .	Db	A9674B ----- 1	
A;Reference number: A86141; MUID:21016719; PMID:11130712	Db	A9674B ----- 1	
A;Accession: A9674B	Db	A9674B ----- 1	
A;Status: preliminary	Db	A9674B ----- 1	
A;Molecule type: DNA	Db	A9674B ----- 1	
A;Residues: 1-946 <STOP>	Db	A9674B ----- 1	
A;Cross-references: UNIPROT:Q9C9D8; GB:AB005173; NID:96730761; PIDN:AAR27150.1; G	Db	A9674B ----- 1	
C;Genetics:	Db	A9674B ----- 1	
A;Gene: TI010.13	Db	A9674B ----- 1	
A;Map position: 1	Db	A9674B ----- 1	
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Score	128.5	Db	128.5
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Length	946;	Db	946;
Mismatches	94;	Db	94;
Conservative	56;	Db	56;
Indels	177;	Db	177;
Gaps	16;	Db	16;
Query Match			
Best Local Similarity	21.0%	Db	21.0%
Score	12.4	Db	12.4
Pred. No.	2.4;	Db	2.4;
Length	946;	Db	946;
Mismatches	94;	Db	94;
Conservative	56;	Db	56;
Indels	121;	Db	121;
Gaps	16;	Db	16;
Query Match			
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Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	61;	Db	61;
Conservative	147;	Db	147;
Indels	138;	Db	138;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
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Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
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Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
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Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
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Score	130;	Db	130;
Pred. No.	5;	Db	5;
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Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
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Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
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Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
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Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
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Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
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Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
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Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
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Length	1927;	Db	1927;
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Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
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Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
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Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative	61;	Db	61;
Indels	147;	Db	147;
Gaps	21;	Db	21;
Query Match			
Best Local Similarity	20.5%	Db	20.5%
Score	130;	Db	130;
Pred. No.	5;	Db	5;
Length	1927;	Db	1927;
Mismatches	69;	Db	69;
Conservative			

Mon Aug 22 13:02:31 2005

us-10-629-951-2.rpr

Oy 383 --DNNCSPTRKDFGEGKIFQEDTIPRT 407
Db : ||| : |:
752 HSGEENCS-----FLPATVPTI 768

Search completed: August 22, 2005, 10:06:00
Job time : 46 secs

GenCore version 5.1.6
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OM protein - protein search, using SW model

Run on: August 22, 2005, 10:02:08 ; Search time 170 Seconds

Scoring table: BL0SUM62

Sequence: 1 MGTTCGLESLSISLGDRGAAPTV. HKLNKVHDMLWENHREKLSLS 580

Post-processing: Minimum Match 0%

Maximum DB seq length: 0

Maximum DB seq length: 200000000

Total number of hits satisfying chosen parameters: 2105692

Minimum DB seq length: 0

Maximum DB seq length: 1319.535 Million cell updates/sec

Scoring table: Gapop 10.0 , Gapext 0.5

Searched: 2105692 seqs, 386760381 residues

Database : A_Geneseq_16Dec04:*

1: geneseqP1980:*

2: geneseqP1990:*

3: geneseqP2000:*

4: geneseqP2019:*

5: geneseqP2002B:*

6: geneseqP2003B:*

7: geneseqP2003B:*

8: geneseqP2004B:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

ALIGNMENTS

Result No.	Score	Query Length	DB ID	Description
1	3055	100.0	580	AAW74473 Human MED
2	3055	100.0	580	Aaw74473 Human MED
3	3055	100.0	580	Add89106 Human 5-m
4	2958.5	96.8	565	Add13954 Human met
5	2194.5	71.8	439	AYY44504 Human del
6	1639	53.6	7	AYY76548 Human ova
7	1230	40.3	257	Add89105 Human 5-m
8	1230	40.3	257	Add89101 Secreted
9	1230	40.3	257	Abb50468 Human sec
10	1230	40.3	257	Abb44725 Novel hum
11	1106	36.2	202	Ab026205 Human pro
12	880.5	28.8	416	Add89108 Chicken 5-m
13	816	26.7	147	Add89117 Human 5-m
14	357	6.8	50	Aae22168 Human MBD
15	254	8.3	4	Abb51147 Human sec
16	254	8.3	50	Ab045404 Novel hum
17	254	8.3	7	Ab026184 Protein a
18	220.5	7.2	467	Adk13968 Frog meth
19	220.5	7.2	467	Adk13967 Frog meth
20	213.5	7.0	486	Adk13972 Human met
21	209	6.8	476	Adk13972 Human hea
22	209	6.8	8	Adk13973 Human met
23	208.5	6.8	477	Adk13961 Human met
24	208.5	6.8	486	Adk13966 Human met
25	208.5	6.8	486	Adk13971 Human met

SUMMARIES

Result ID	Date	Description
AAW74473	19-MAY-1999	(first entry)
XX	DE	Human MED1 endonuclease protein sequence.
XX	KW	Endonuclease; MED1; human; methyl-CPG binding endonuclease-1; DNA fidelity; DNA manipulation; cancer; fragile X syndrome; therapy; myotonic dystrophy; Huntington's disease; spinocerebellar ataxia; Kennedy's disease; triplet repeat expansion disorder.
XX	OS	Homo sapiens.
XX	PN	WO9904626-A1.
XX	PD	04-FEB-1999.
XX	PR	28-JUL-1998; 98WO-US0150828.
XX	PR	28-DECEMBER-1997; 97US-0053936P.
XX	PA	(FOXC-) FOX CHASE CANCER CENT.
XX	PI	Bellacosa A;
XX	DR	WPI: 1999-142462/12.
XX	DR	N-PSDB; AxX22002.
XX	PT	New nucleic acid encoding human endonuclease MED1 involved in DNA mismatch repair - used for diagnosing susceptibility to cancer and fragile X syndrome, and therapeutically.
XX	PS	Claim 8; Fig 3; 109pp; English.

This sequence is the human MED1 endonuclease of the invention. MED1 (for methyl-CPG binding endonuclease-1) is used to screen for specific modulators (potential therapeutic agents particularly mimetics of MED1) and to study interactions involved in maintaining DNA fidelity, for DNA manipulation and to raise antibodies. Susceptibility or predisposition to cancer (particularly colorectal cancer), or its prognosis, where caused by -polypsis colorectal cancer), or its prognosis, where caused by sequence alterations in the MED1-encoding gene, are identified by sequence comparison, amplification, detection, altered polypeptide, and restriction fragment mapping, hybridisation (particularly to probes specific for a

CC mutant allele). Those same methods can also be used to diagnose fragile X syndrome and other diseases (e.g., myotonic dystrophy, Huntington's disease, spinocerebellar ataxia and Kennedy's disease) associated with triplet repeat expansion. The DNA, or its fragments, are used as probes and primers in the above diagnostic methods, also to isolate homologous sequences, as sources of antisense sequences and for gene transfer, particularly to restore drug sensitivity to drug-resistant cancer cells.

XX SQ Sequence 580 AA;

Query Match 100.0%; Score 3055; DB 2; Length 580;
Best Local Similarity 100.0%; Pred No. 7.7e-287; Length 580;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CNPLIQLQPTIASAQFGATAGTECRKSVP CGWERTVKQRLFGKTAGFDVYFISPOGLKERS 60
Db 1 MGTTGLESISLGDRGAAPVTSSRLVDPDNDLKEVDAMELERVGDEBEMMIKRSSE 60

Qy 61 CNPLIQLQPTIASAQFGATAGTECRKSVP CGWERTVKQRLFGKTAGFDVYFISPOGLKERS 60
61 ONPLIQEPIASAQFGATAGTECRKSVP CGWERTVKQRLFGKTAGFDVYFISPOGLKERS 120

Qy 121 KSSLANYLHKGETSLKPEDDFPDLVSKRGKSRVYKDCSMAALTSHLQNOQNNSNNWLR 180
121 KSSLANYLHKGETSLKPEDDFPDLVSKRGKSRVYKDCSMAALTSHLQNOQNNSNNWLR 180

Db 181 RSKCKKDVFMPSSSELOPSRGLNPTSTHLLKEDEGTDVNFRKVPKPKVTLKG 240
181 RSKCKKDVFMPSSSELOPSRGLNPTSTHLLKEDEGTDVNFRKVPKPKVTLKG 240

Qy 241 IPIKKTKGCRKSCSGFWQDSKRSVNCDAESEPQAKSOLRTVTCISDAGCETL 300
241 IPIKKTKGCRKSCSGFWQDSKRSVNCDAESEPQAKSOLRTVTCISDAGCETL 300

• Qy 301 SVTSBENSLVKRKERSLSGSNFCSBQKTSGGINIFCSPACKSERNEKEDTFLESBEIGT 360
301 SVTSBENSLVKRKERSLSGSNFCSBQKTSGGINIFCSPACKSERNEKEDTFLESBEIGT 360

Qy 361 KVEVERKEHLHTDILKGSBMDNCSPTRKDFTGEKIFOEDTIPRTQIERKTSLYFSS 420
361 KVEVERKEHLHTDILKGSBMDNCSPTRKDFTGEKIFOEDTIPRTQIERKTSLYFSS 420

Db 361 KVEVERKEHLHTDILKGSBMDNCSPTRKDFTGEKIFOEDTIPRTQIERKTSLYFSS 420

RESULT 2

ADD89906 ID ADD89906 standard; protein; 580 AA.

AC ADD89906; DT 29-JAN-2004 (first entry)

DE Human 5-methylcytosine DNA glycosylase.

KW Human; 5-methylcytosine DNA glycosylase; enzyme; Cpg.

XX OS Homo sapiens.

XX PN WO2003078593-A2.

XX PD 25-SEP-2003.

PF 14-MAR-2003; 2003WO-US007933.

XX PR 15-MAR-2002; 2002US-036468P.

XX PA (EPIC-) EPIGENOMICS AG.

XX PI Lofton-Day CE, Day JK.

XX DR WPI; 2003-779127/73.

DR N-PSDB; ADD89905.

XX PT Labeling methylated or methylatable CpG sequences, useful e.g. for diagnostic detection of altered methylation, comprises replacing methylated cytosine by labeled cytosine.

XX PS Claim 11; Page 53-55; 73pp; English.

CC The present sequence is the protein sequence of human 5-methylcytosine DNA glycosylase (5-McDG). The enzyme acts by cleaving glycosylic bonds at methylated CpG sites of DNA, removing 5-methylcytosine from the DNA backbone as a free base. Human 5-McDG can be used in a claimed method for labeling CpG sequences corresponding to methylated CpG sequences in an isolated DNA sample. The method comprises: digesting the genomic DNA with a restriction endonuclease to produce genomic DNA fragments; treating the genomic DNA fragments with 5-McDG such that one or more 5-methylcytosine bases are removed to produce abasic genomic DNA fragments; and treating these abasic genomic DNA fragments with base excision repair enzymes in the presence of labelled dcmP such that 5-methylcytosine removed from the genomic DNA fragments by 5-McDG is replaced by labelled cytosine in the one or more corresponding positions of the abasic genomic DNA fragments to produce labelled genomic DNA fragments, so that specific labelling of CpG sequences corresponding to methylated CpG sequences is achieved. The 5-McDG is also used in a claimed method for comparing CpG methylation status, extent or pattern between or among reference and test genomic DNA samples, and in a claimed method for labelling potentially-methylatable CpG sequences in CpG-containing genomic DNA fragments. The methods are used to identify methylated and/or potentially methylatable CpG dinucleotides in genomic DNA, including comparison of methylation pattern between healthy and diseased samples, for diagnosis.

XX SQ Sequence 580 AA;

Query Match 100.0%; Score 3055; DB 7; Length 580;
Best Local Similarity 100.0%; Pred No. 7.7e-287; Length 580;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MGTTGLESISLGDRGAAPVTSSRLVDPDNDLKEVDAMELERVGDEBEMMIKRSSE 60
1 MGTTGLESISLGDRGAAPVTSSRLVDPDNDLKEVDAMELERVGDEBEMMIKRSSE 60

Db 181 RSKCKKDVFMPSSSELOPSRGLNPTSTHLLKEDEGTDVNFRKVPKPKVTLKG 240
181 RSKCKKDVFMPSSSELOPSRGLNPTSTHLLKEDEGTDVNFRKVPKPKVTLKG 240

Qy 241 IPIKKTKGCRKSCSGFWQDSKRSVNCDAESEPQAKSOLRTVTCISDAGCETL 300
241 IPIKKTKGCRKSCSGFWQDSKRSVNCDAESEPQAKSOLRTVTCISDAGCETL 300

Db 241 IPIKKTKGCRKSCSGFWQDSKRSVNCDAESEPQAKSOLRTVTCISDAGCETL 300

Qy 301 SVTSBENSLVKRKERSLSGSNFCSBQKTSGGINIFCSPACKSERNEKEDTFLESBEIGT 360
301 SVTSBENSLVKRKERSLSGSNFCSBQKTSGGINIFCSPACKSERNEKEDTFLESBEIGT 360

Qy 361 KVEVERKEHLHTDILKGSBMDNCSPTRKDFTGEKIFOEDTIPRTQIERKTSLYFSS 420
361 KVEVERKEHLHTDILKGSBMDNCSPTRKDFTGEKIFOEDTIPRTQIERKTSLYFSS 420

QY 421 KYNKEALSPPRKAFKKWTPRSPFNLVQETLFHDPMKLLATIPLRTSGKMAIPVLWK 480
Db 421 KYNKEALSPPRKAFKKWTPRSPFNLVQETLFHDPMKLLATIPLRTSGKMAIPVLWK 480
QY 481 FLEKYPSAEVARTADWRDVSLILKPGLYDRAKTIVKFSDEYLTKWKPIELHIGKY 540
Db 481 FLEKYPSAEVARTADWRDVSLILKPGLYDRAKTIVKFSDEYLTKWKPIELHIGKY 540
QY 541 GNDSYRIFCNEWKQVHPEDIKLNKTHDWLWNHENHEKLSSL 580
Db 541 GNDSYRIFCNEWKQVHPEDIKLNKTHDWLWNHENHEKLSSL 580

RESULT 3
ID ADKL3954 standard; protein; 580 AA.
XX ADKL3954;
XX DT 03-JUN-2004 (First entry)
XX DE Human methyl-CpG-binding protein #9.
XX KW Rett syndrome; methyl-CpG-binding protein 2; MECP2;
KW neurodevelopmental disease; autism; non-syndromic mental retardation;
KW idiopathic neonatal encephalopathy; idiopathic infantile spasm;
KW idiopathic cerebral palsy; Angelman syndrome; schizophrenia; human.
OS Homo sapiens.
XX PN US6709817-B1.
XX PD 23-MAR-2004.
XX PP 07-SEP-2000; 2000US-00657013.
PR 07-SEP-1999; 99US-0152778P.
XX PA (BAYU) BAYLOR COLLEGE MEDICINE.
XX PI Zoghbi HY, Van den Veyver IB, Amir R, Francke U;
XX DR WPI; 2004-256068/24.
XX PT Screening human for Rett syndrome comprises detecting mutation in nucleic acid sequence encoding methyl-CpG-binding protein 2 (MECP2).
XX PS Disclosure; SEQ ID NO 56; 125pp; English.
XX CC The invention relates to a method of screening a human for Rett syndrome comprising detecting a mutation in a nucleic acid sequence encoding methyl-CpG-binding protein 2 (MECP2). The method is useful for screening a human for Rett syndrome. The method is useful for screening neurodevelopmental diseases such as Rett syndrome, autism, non-syndromic mental retardation, idiopathic neonatal encephalopathy, idiopathic infantile spasms, idiopathic cerebral palsy, Angelman syndrome and schizophrenia. The present sequence represents the amino acid sequence of a methyl-CpG-binding protein.
XX SQ Sequence 580 AA;

Query Match 100.0%; Score 3055; DB 8; length 580;
Best Local Similarity 100.0%; Pred. No. 7; re-207;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MGTRGLESILSLGGRGAAPPTVSSRLVDPDPLRKERYAVAMELERVGDBEOMNIKRSSE 60
Db 1 MGTRGLESILSLGGRGAAPPTVSSRLVDPDPLRKERYAVAMELERVGDBEOMNIKRSSE 60

QY 61 CNPLQEPITASAQGATAGTCTCRRSVPQWGRWVQRLFGKTAGRFDYFISPOGLKRS 120
Db 61 CNPLQEPITASAQGATAGTCTCRRSVPQWGRWVQRLFGKTAGRFDYFISPOGLKRS 120

QY 121 KSSLANYLHKNGETSLKPBDPFDFTVLSKGKRSYKDCSNALTSHLQNSNNSNWLRT 180
Db 121 KSSLANYLHKNGETSLKPBDPFDFTVLSKGKRSYKDCSNALTSHLQNSNNSNWLRT 180
QY 181 RSKCKDWFMPSSSELQSRGLSNFTSHLILKEDGVDDVNRKVRKPKGKTYLKG 240
Db 181 RSKCKDWFMPSSSELQSRGLSNFTSHLILKEDGVDDVNRKVRKPKGKTYLKG 240
QY 241 IPIKTKKGCRKSCGGFVQDSKRSVNCADALESEPVAKSOLDRTVCTSDAGACETL 300
Db 241 IPIKTKKGCRKSCGGFVQDSKRSVNCADALESEPVAKSOLRTVCTSDAGACETL 300
QY 301 SVTSEENSLVKKERSSNSFCBOKTSQGIIINKPCSAKOSERHNBEYDTFLESBEGT 360
Db 301 SVTSEENSLVKKERSSNSFCBOKTSQGIIINKPCSAKOSERHNBEYDTFLESBEGT 360
QY 361 KVEWVERKERHLHTDILKGSMDNINCPTKDFGKIRDKTIRTQIRRKTSLYFSS 420
Db 361 KVEWVERKERHLHTDILKGSMDNINCPTKDFGKIRDKTIRTQIRRKTSLYFSS 420
QY 421 KYNKEALSPPRKAFKKWTPRSPFNLVQETLFHDPMKLLATIPLRTSGKMAIPVLWK 480
Db 421 KYNKEALSPPRKAFKKWTPRSPFNLVQETLFHDPMKLLATIPLRTSGKMAIPVLWK 480
QY 481 FLEKYPSAEVARTADWRDVSLILKPGLYDRAKTIVKFSDEYLTKWKPIELHIGKY 540
Db 481 FLEKYPSAEVARTADWRDVSLILKPGLYDRAKTIVKFSDEYLTKWKPIELHIGKY 540
QY 541 GNDSYRIFCNEWKQVHPEDIKLNKTHDWLWNHENHEKLSSL 580
Db 541 GNDSYRIFCNEWKQVHPEDIKLNKTHDWLWNHENHEKLSSL 580

RESULT 4
ID AAY44504 standard; protein; 565 AA.
XX AAY44504;
XX AC AAY44504;
XX DT 27-MAR-2000 (First entry)
XX DE Human delta228-UV damage endonuclease.
XX KW Delta228-UVDE; ultraviolet damage endonuclease; GST signal peptide;
KW glutathione-S-transferase; signal peptide; uvc1+ gene product;
KW UV irradiation; DNA damage; UV radiation damage; photoproduct;
KW abasic site; apatinium dianuct; mismatched nucleotide pairing;
KW nucleotide alkylation; skin cancer.

XX OS Homo sapiens.
XX PN W09963828-A1.
XX PD 16-DEC-1999.
XX PF 08-JUN-1999; 99WO-US012910.
XX PR 08-JUN-1998; 99US-008851P.
PR 18-MAY-1999; 99US-013472P.
XX PA (UYEM-) UNIV EMORY.
XX PI Doetsch PW, Kaur B, Avery AM;
DR WPI; 2000-116417/10.
XX PT A new truncated ultraviolet damage endonuclease for treatment of skin cancers.
XX PS Claim 16; Page 60; 133pp; English.
CC The present sequence is human delta228-UV damage endonuclease. Delta228-

Db	413	KTIVKFSDEYLTKQWKPPIELHIGIG	437		Db	1	ESEPVAAQSKQLDRTWCISAGACETLSVTSEENSLVKKERSLSSGSNFCSEQKTSGII	60	
RESULT 6					Db	334	NKFCSAKDSHENEKEDTFLESBEGTKEVERGEHLHDILRGSEMDDNCSPTRKDF	393	
ADD8915	ID	ADD8915 standard; protein;	307 AA.		Db	61	NKFCSAKDSHENEKEDTFLESBEGTKEVERGEHLHDILRGSEMDDNCSPTRKDF	120	
XX	XX				Db	394	TGECIFQODDTIPRTQERRKTSLYSSKYNKEALSPPRRCAFPKKTPPRSPFNLVQETLF	453	
AC	AC	ADD8915;			Db	121	TGECIFQODDTIPRTQERRKTSLYSSKYNKEALSPPRRCAFPKKTPPRSPFNLVQETLF	180	
DT	29-JAN-2004	(first entry)			Db	454	HDPWKLLIATIFLNRTSKGKMAIPVWKFLLKYPSSAVARTADWRDVSELKPLGLYDLRA	513	
XX	XX	Human 5-methylcytosine DNA glycosylase N-terminal deletion mutant.			Db	181	HDPWKLLIATIFLNRTSKGKMAIPVWKFLLKYPSSAVARTADWRDVSELKPLGLYDLRA	240	
KW	XX	Human; 5-methylcytosine DNA glycosylase; enzyme; Cpg; mutant; mutein.			Db	514	KTIVKFSDEYLTKQWKPPIELHIGIGKGNDSYRITCWNEMQVHPEDHKUNYHWLWEN	573	
OS	XX	Homo sapiens.			Db	241	KTIVKFSDEYLTKQWKPPIELHIGIGKGNDSYRIFCVNEWQVHPEDHKUNYHWLWEN	300	
PV	PN	WO2003078593-A2.			Db	QY	574	HBKLISL 580	
XX	XX	25-SEP-2003.			Db	QY	301	HBKLISL 307	
PP	PT	14-MAR-2003; 2003WO-US007933.			RESULT 7				
XX	XX	15-MAR-2002; 2002US-0364689P.			Db	AAW88701			
XX	PA	(EPIG-) EPIGENOMICS AG.			Db	AAW88701			
XX	PT	Lofton-Day CB, Day JK;			Db	AAW88701;			
XX	DR	WPI; 2003-779127/73.			Db	AAW88701;			
XX	PT	Labeling methylated or methylatable Cpg sequences, useful e.g. for diagnostic detection of altered methylation, comprises replacing methylated cytosine by labeled cytosine.			Db	01-MAR-1999	(first entry)		
XX	PT	The present sequence is the protein sequence of an N-terminal deletion mutant of human 5-methylcytosine DNA glycosylase (5-MCDG), in which amino acid residue 1 corresponds to amino acid 274 of the full-length protein ADD8906. 5-MCDG acts by cleaving glycosidic bonds at methylated Cpg sites of DNA, removing 5-methylcytosine from the DNA backbone as a free base. The N-terminal deletion mutant shows enhanced deglycosylase specificity towards Cpg dinucleotide sequences. Human 5-MCDG can be used in a claimed method for labelling Cpg sequences corresponding to methylated Cpg sequences in an isolated DNA sample. The method comprises: digesting the genomic DNA with a restriction endonuclease to produce genomic DNA fragments; treating the genomic DNA fragments with 5-MCDG such that one or more 5-methylcytosine bases are removed to produce abasic genomic DNA fragments; and treating those abasic genomic DNA fragments with base excision repair enzymes in the presence of labelled dCTP such that 5-methylcytosine removed from the genomic DNA fragments by 5-MCDG is replaced by labelled cytosine in the one or more corresponding positions of the abasic genomic DNA fragments to produce labelled genomic DNA fragments, so that specific labelling of Cpg sequences corresponding to methylated Cpg sequences is achieved. The 5-MCDG is also used in a claimed method for comparing Cpg methylation status, extent or pattern between or among reference and test genomic DNA samples, and in a claimed method for labelling potentially-methylatable Cpg sequences in Cpg-containing genomic DNA fragments. The methods are used to identify methylated and/or potentially methylatable Cpg dinucleotides in genomic DNA, including comparison of methylation pattern between healthy and diseased samples, for diagnosis.			Db	04-JUN-1998;	98WO-US011422.		
XX	CC	Sequence 307 AA;			Db	06-JUN-1997;	97US-00489375P.		
CC	CC	Query Match Similarity 53.6%; Score 1639; DB 7; Length 307; Best Local Similarity 100.0%; Pred. No. 8; 4e-150; Mismatches 0; Indels 0; Gaps 0;			Db	06-JUN-1997;	97US-0048881P.		
CC	CC	Matches 307; Conservative 0; MisMatches 0; Indels 0; Gaps 0;			Db	06-JUN-1997;	97US-0048882P.		
CC	CC	274 ESPVVAQSKQLDRTWCISAGACETLSVTSEENSLVKKERSLSSGSNFCSEQKTSGII 333			Db	06-JUN-1997;	97US-0048915P.		
CC	CC	06-JUN-1997;			Db	06-JUN-1997;	97US-0048897P.		
XX	SQ	Sequence 307 AA;			Db	06-JUN-1997;	97US-0048888P.		
XX	XX	Query Match Similarity 53.6%; Score 1639; DB 7; Length 307; Best Local Similarity 100.0%; Pred. No. 8; 4e-150; Mismatches 0; Indels 0; Gaps 0;			Db	06-JUN-1997;	97US-0048899P.		
CC	CC	Matches 307; Conservative 0; MisMatches 0; Indels 0; Gaps 0;			Db	06-JUN-1997;	97US-0048901P.		
CC	CC	06-JUN-1997;			Db	06-JUN-1997;	97US-0048917P.		

PR 06-JUN-1997; 97US-0048949P.
 PR 06-JUN-1997; 97US-0048962P.
 PR 06-JUN-1997; 97US-0048963P.
 PR 06-JUN-1997; 97US-0048964P.
 PR 06-JUN-1997; 97US-0048970P.
 PR 06-JUN-1997; 97US-0048971P.
 PR 06-JUN-1997; 97US-0048972P.
 PR 06-JUN-1997; 97US-0048974P.
 PR 06-JUN-1997; 97US-0049019P.
 PR 06-JUN-1997; 97US-0049020P.
 PR 06-JUN-1997; 97US-0049373P.
 PR 06-JUN-1997; 97US-0049374P.
 PR 06-JUN-1997; 97US-0049375P.
 PR 05-SEP-1997; 97US-0057584P.
 PR 05-SEP-1997; 97US-0057627P.
 PR 05-SEP-1997; 97US-0057628P.
 PR 05-SEP-1997; 97US-0057629P.
 PR 05-SEP-1997; 97US-0057634P.
 PR 05-SEP-1997; 97US-0057635P.
 PR 05-SEP-1997; 97US-0057642P.
 PR 05-SEP-1997; 97US-0057643P.
 PR 05-SEP-1997; 97US-0057644P.
 PR 05-SEP-1997; 97US-0057646P.
 PR 05-SEP-1997; 97US-0057647P.
 PR 05-SEP-1997; 97US-0057648P.
 PR 05-SEP-1997; 97US-0057649P.
 PR 05-SEP-1997; 97US-0057650P.
 PR 05-SEP-1997; 97US-0057651P.
 PR 05-SEP-1997; 97US-0057654P.
 PR 05-SEP-1997; 97US-0057661P.
 PR 05-SEP-1997; 97US-0057662P.
 PR 05-SEP-1997; 97US-0057666P.
 PR 05-SEP-1997; 97US-0057667P.
 PR 05-SEP-1997; 97US-0057668P.
 PR 05-SEP-1997; 97US-0057760P.
 PR 05-SEP-1997; 97US-0057761P.
 PR 05-SEP-1997; 97US-0057762P.
 PR 05-SEP-1997; 97US-0057763P.
 PR 05-SEP-1997; 97US-0057764P.
 PR 05-SEP-1997; 97US-0057765P.
 PR 05-SEP-1997; 97US-0057769P.
 PR 05-SEP-1997; 97US-0057770P.
 PR 05-SEP-1997; 97US-0057771P.
 PR 05-SEP-1997; 97US-0057774P.
 PR 05-SEP-1997; 97US-0057775P.
 PR 05-SEP-1997; 97US-0057776P.
 PR 05-SEP-1997; 97US-0057777P.
 PR 05-SEP-1997; 97US-0057778P.
 PR 18-DEC-1997; 97US-0070923P.

Sequence 257 AA;
 Query Match Score 1230; DB 2: Length 257;

Best local similarity 93.1%; Pred. No. 3..3e-110; Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;

QY	160	MAAITSHQNQSNNSNNWTRTSKCKQVFMPPSSPLQESRGLSNTSTHILKEPG	219
Db	1	MAAITSHQNQSNNSNNWTRTSKCKQVFMPPSSPLQESRGLSNTSTHILKEPG	60
QY	220	VDDVNPKRKPKPKGKVTLKGPIKKTKGCRKSCSGVQDSKRESYCNKADAESPV	279
Db	61	VDDVNPKRKPKPKGKVTLKGPIKKTKGCRKSCSGVQDSKRESYCNKADAESPV	120
QY	280	QKSOLDRTWCISDAGACSETLSTVSEENSLVKCKERISISSLGSNPFCEORTGJINKFGSA	339
Db	121	QKSOLDRTWCISDAGACSETLSTVSEENSLVKCKERISISSLGSNPFCEORTGJINKFGSA	180
QY	340	KDSHNEKYEEDTPELESEBEGTKYEVVERKEHLHTDLRGSEMDNNNSPDKFTGKTF	399
Db	181	KDSHNEKYEEDTPELESEBEGTKYEVVERKEHLHTDLRGSEMDNNNSPDKFT-----	235
QY	400	QEDTIPRQIERRTKLSLP	418
Db	236	-EDTIPRQIDRKKENPKVF	253

RESULT 8
 ABB50468
 ID ABB50468 standard; protein; 257 AA.
 XX
 AC ABB50468;
 XX DT 07-FEB-2002 (first entry)
 DE Human secreted protein encoded by gene 168 SEQ ID NO:416.
 XX Human; secreted protein; immunomodulatory; anti-sclerotic; anti-HIV;
 KW dermato logical; immunosuppressive; antiinflammatory; immunostimulant;
 KW cyrostatic; cardiant; vascular; antiangiogenic; opthalmological;
 KW neuroprotective; nootropic; anticonvulsant; anti-alzheimer's; vulnery;
 KW anti-parkinsonian; antimicrobial; gene therapy; vaccine; immune disorder;
 KW multiple sclerosis; systemic lupus erythematosus; HIV infection; cancer;
 KW human immunodeficiency virus; hyperproliferative disorder; wound healing;
 KW Gaucher's disease; cardiovascular disease; scimitar syndrome; chemotaxis;
 KW Chaga's cardiomyopathy; coronary arteriosclerosis; angiogenic disorder;
 KW corneal graft neovascularization; diabetic retinopathy; regeneration;
 KW neurological disorder; Huntington's chorea; Alzheimer's disease;
 KW Parkinson's disease; infectious disease; chromosome 3.
 XX OS Homo sapiens.

(HUMA-) HUMAN GENOME SCI INC.

PI Young P, Greene JM, Ferrie AM, Ruben SM, Rosen CA, Hu J;
 PI Olsen HS, Ebner R, Brewer LA, Moore PA, Shi Y, Florence C;
 PI Florence K, Lafleur DW, Ni J, Pan P, Wei Y, Fischer CL, Soppet DR;
 PI Li Y, Zeng Z, Kyaw H, Yu G, Feng P, Dillon PJ, Endress GA;
 PI Carter KC;
 XX DR WPI; 1999-059865/05.
 DR N-PSDB; AAV84578.

XX New isolated human genes and the secreted polypeptides they encode - useful for diagnosis and treatment of e.g. cancers, neurological disorders, immune diseases, inflammation or blood disorders.

XX The invention relates to nucleic acid sequences (AAV8411 to AAV84633) encoding human secreted proteins (AAW8534 to AAW88756). The secreted protein gene sequences are deposited with the ATCC under deposit numbers ATCC 9799, 9794, 9795, 9796, 9797, 209009, 209010, 209011, 209012, 209013, 209014, 209015, 209016, 209017, 209018, 209019, 209020, 209021, 209022, 209023, 209024, 209025, 209026, 209027, 209028, 209029, 209030, 209031, 209032, 209033, 209034, 209035, 209036, 209037, 209038, 209039, 209040, 209041, 209042, 209043, 209044, 209045, 209046, 209047, 209048, 209049, 209050, 209051, Host CC

PN WO200162891-A2.
 XX
 PD 30-AUG-2001.
 XX
 PR 21-FEB-2001; 2001WO-US005614.
 XX
 PR 24-FEB-2000; 2000US-0184836P.
 XX
 PR 29-MAR-2000; 2000US-0193170P.
 XX
 PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 NI J, Ebner R, Lafleur DW, Moore PA, Olsen HS, Rosen CR,
 PI Ruben SM, Soppet DR, Young PE, Shi Y, Florence KA, Wei Y,
 PI Florence C, Hu J, Li Y, Kyaw H, Fischer CL, Ferrie AM, Fan P,
 PI Feng P, Endress GA, Dillon PJ, Carter KC, Brewer LA, Yu G, Zeng Z,
 PI Greene JM;
 XX
 WPI; 2001-62574/72.
 DR N-PSDB; ABA83361.
 XX
 PT Nucleic acids encoding 207 human secreted polypeptides, useful for
 PT preventing, diagnosing and/or treating, e.g. cancers, Parkinson's disease
 PT and diabetic retinopathy.
 XX
 PS Claim 11; Page 1171-1172; 1533pp; English.
 XX
 CC ABB50301 to ABB51287 and ABA83194 to ABA83441 represent human secreted
 CC proteins (I) and poly-nucleotide (II) sequences. (I) and (II) have various
 CC activities based on the tissues and cells the genes are expressed in.
 CC Example of these activities include: immunomodulatory; anti-angiogenic;
 CC dermatological; immunosuppressive; anti-inflammatory; ophthalmological;
 CC anti-HIV; cytostatic; cardiotonic; anti-angiogenic; opthalmological;
 CC neuroprotective; noctropic; anticonvulsant; antialzheimers; vascular;
 CC anti-parkinsonian; antimicrobial; and vulnerary. (I) and (II) can be used
 CC in gene therapy and vaccine production. (I) and (II) can be used in the
 CC prevention, diagnosis and treatment of immune disorders (e.g. multiple
 CC sclerosis, systemic lupus erythematosus and human immunodeficiency virus
 (HIV) infections), hyperproliferative disorders (e.g. cancers and
 Gaucher's disease), cardiovascular diseases (e.g. Scimitar Syndrome,
 Chagot's cardiomyopathy and coronary arteriosclerosis), angiogenic
 CC disorders (e.g. corneal graft neovascularisation and diabetic
 CC retinopathy), neurological disorders (e.g. Huntington's chorea,
 CC Alzheimer's disease and Parkinson's disease), infectious diseases and/or
 CC for promoting wound healing, regeneration and/or chemotaxis. ABA83185 to
 CC ABA83193 and ABB50300 represent sequences used in the exemplification of
 CC the present invention
 XX
 SQ Sequence 257 AA;

Query Match 40.3%; Score 1230; DB 4; Length 257;
 Best Local Similarity 93.1%; Pred. No. 3; 3e-110;
 Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;

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  QY 160 MAALTSILQNGNSNNWNLRSCKKDVMPFMPSSSEBLOESRGLNFSTHLLKEDEC 219
      ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
  Db 1 MAALTSHLQNQNSNNWNLRSCKKDVMPFMPSSSEBLOESRGLNFSTHLLKEDEC 60
      ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
  QY 220 VDDVNFRKVKGPKGVTLKGIPKKRKGRKCSGCFVQSDSKRSEVNCNADAESEPV 279
      ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
  Db 61 VDDVNFRKVKGPKGVTLKGIPKKRKGRKCSGCFVSDSKRSEVNCNADAESEPVA 120
      ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
  QY 340 KOSEHNKEPTEPLESIEIGTKEVERKEHHTDILKGSMDNCSPTKDFTRKIF 399
      ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
  Db 181 KOSEHNKEPTEPLESIEIGTKEVERKEHHTDILKGSMDNCSPTKDFTRKIF----- 235
      ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
  QY 400 QEDTIPRTQIERKRTSLYF 418
      ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
  Db 236 -EDTIPRTQIERKRTSLYF 253
      ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
  
```

RESULT 9
 ABO4725
 ID ABO4725 standard; protein; 257 AA.
 XX
 AC ABO4725;
 XX
 DT 02-OCT-2003 (first entry)
 DE Novel human secreted protein #168.
 XX
 Human; gene therapy; autoimmune disorder; multiple sclerosis; cancer;
 KW systemic lupus erythematosus; haemopoietic cell disorder; allergy;
 KW agammaglobulinaemia; ataxia telangiectasia; blood coagulation disorder;
 KW fibrinogenemia; thrombocytopenia; graft-versus-host disease; arthritis;
 KW inflammatory condition; ischaemia-reperfusion injury; infectious disease;
 KW hyperproliferative disorder; purpura; viral infection; regeneration;
 KW bacterial infection; ulcer; Alzheimer's disease.
 XX
 OS Homo sapiens.
 XX
 PN US2003065160-A1.
 XX
 PD 03-APR-2003.
 XX
 DP 07-DEC-2001; 2001US-00004860.
 XX
 PR 06-JUN-1997; 97US-0048875P.
 PR 06-JUN-1997; 97US-0048876P.
 PR 06-JUN-1997; 97US-0048877P.
 PR 06-JUN-1997; 97US-0048878P.
 PR 06-JUN-1997; 97US-0048880P.
 PR 06-JUN-1997; 97US-0048881P.
 PR 06-JUN-1997; 97US-0048882P.
 PR 06-JUN-1997; 97US-0048883P.
 PR 06-JUN-1997; 97US-0048884P.
 PR 06-JUN-1997; 97US-0048885P.
 PR 06-JUN-1997; 97US-0048892P.
 PR 06-JUN-1997; 97US-0048893P.
 PR 06-JUN-1997; 97US-0048894P.
 PR 06-JUN-1997; 97US-0048895P.
 PR 06-JUN-1997; 97US-0048896P.
 PR 06-JUN-1997; 97US-0048897P.
 PR 06-JUN-1997; 97US-0048898P.
 PR 06-JUN-1997; 97US-0048899P.
 PR 06-JUN-1997; 97US-0048900P.
 PR 06-JUN-1997; 97US-0048901P.
 PR 06-JUN-1997; 97US-0048916P.
 PR 06-JUN-1997; 97US-0048917P.
 PR 06-JUN-1997; 97US-0048949P.
 PR 06-JUN-1997; 97US-0048862P.
 PR 06-JUN-1997; 97US-0048863P.
 PR 06-JUN-1997; 97US-0048864P.
 PR 06-JUN-1997; 97US-0048870P.
 PR 06-JUN-1997; 97US-0048871P.
 PR 06-JUN-1997; 97US-0048872P.
 PR 06-JUN-1997; 97US-0048873P.
 PR 06-JUN-1997; 97US-0048874P.
 PR 06-JUN-1997; 97US-0048875P.
 PR 06-JUN-1997; 97US-0048876P.
 PR 06-JUN-1997; 97US-0048877P.
 PR 06-JUN-1997; 97US-0048878P.
 PR 06-JUN-1997; 97US-0048880P.
 PR 06-JUN-1997; 97US-0048881P.
 PR 06-JUN-1997; 97US-0048882P.
 PR 06-JUN-1997; 97US-0048883P.
 PR 06-JUN-1997; 97US-0048884P.
 PR 06-JUN-1997; 97US-0048885P.
 PR 06-JUN-1997; 97US-0048892P.
 PR 06-JUN-1997; 97US-0048893P.
 PR 06-JUN-1997; 97US-0048894P.
 PR 06-JUN-1997; 97US-0048895P.
 PR 06-JUN-1997; 97US-0048896P.
 PR 06-JUN-1997; 97US-0048897P.
 PR 06-JUN-1997; 97US-0048898P.
 PR 06-JUN-1997; 97US-0048899P.
 PR 06-JUN-1997; 97US-0048900P.
 PR 06-JUN-1997; 97US-0048901P.

PR	05-SEP-1997;	97US-0057646P.	OY	220 VDDVNFRKVRPKKGKVTLKGPIKKTRGCRKCSGIVQSDSKRESYCNKADAESEPVVA 279
PR	05-SEP-1997;	97US-0057647P.	Db	61 VDDVNFRKVRPKKGKVTLKGPIKKTRGCRKCSGIVQSDSKRESYCNKADAESEPVVA 120
PR	05-SEP-1997;	97US-0057648P.	OY	280 QKSOLDRTVCISDAGACCTLSTPSEEENSLVCKERSSISGSNFCSEOKTSGINKFCSA 339
PR	05-SEP-1997;	97US-0057650P.	Db	121 QKSOLDRTVCISDAGACCTLSTPSEEENSLVCKERSSISGSNFCSEOKTSGINKFCSA 180
PR	05-SEP-1997;	97US-0057651P.	OY	340 KDSHENEYKEDTELESEGTKEVVERKEHAIHTDLRGSEMDNNASPTKDFTGEKL 399
PR	05-SEP-1997;	97US-0057662P.	Db	181 KDSHENEYKEDTELESEGTKEVVERKEHAIHTDLRGSEMDNNASPTKDFT----- 235
PR	05-SEP-1997;	97US-0057667P.	OY	400 QEDTIPRQIERRKTSLYF 418
PR	05-SEP-1997;	97US-0057668P.	Db	236 -EDTIPRQIERRKTSLYF 253
PR	05-SEP-1997;	97US-0057765P.		
PR	05-SEP-1997;	97US-0057769P.		
PR	05-SEP-1997;	97US-0057770P.		
PR	05-SEP-1997;	97US-0057771P.		
PR	05-SEP-1997;	97US-0057774P.		
PR	05-SEP-1997;	97US-0057775P.		
PR	05-SEP-1997;	97US-0057776P.		
PR	05-SEP-1997;	97US-0057777P.		
PR	05-SEP-1997;	97US-0057778P.		
PR	04-JUN-1998;	98US-0011422.		
PR	15-JUL-1998;	98US-0092921P.		
PR	30-JUL-1998;	98US-0092921P.		
PR	04-DEC-1998;	98US-00205258.		
PA	(HUMA-) HUMAN GENOME SCI INC.			
XX			RESULT 10	
XX			ABO26205	
XX			ID ABO26205 standard; protein; 257 AA.	
XX			AC ABO26205;	
XX			XX	
DT	10-SEP-2003	(first entry)		
XX			DE Human protein from novel secreted protein gene 168.	
XX			KW Human; secreted protein; precerebellin-like protein;	
XX			KW neurodegenerative disorder; behavioural disorder; Alzheimer's disease;	
XX			KW Parkinson's disease; Huntington's disease; schizophrenia; mania;	
XX			KW dementia; paranoia; psychosis; autism; immune disorder; infection;	
XX			KW inflammation; allergy; liver disorder; hepatoblastoma; Jaundice;	
XX			KW hepatitis; immunological disorder; AIDS; leukaemia; rheumatoid arthritis;	
XX			KW sepsis; acne; psoriasis; cancer.	
OS			XW Homo sapiens.	
XX			PN US6525174-B1.	
XX			PD 25-FEB-2003.	
XX			PP 04-DEC-1998;	
PT	New isolated protein, useful for preparing a composition for diagnosing	98US-00205258.	PP	
PT	or treating cancer, inflammatory, immune or infectious diseases.		XX	
XX			PR 06-JUN-1997;	
PS	Disclosure; SEQ ID NO 416; 172pp; English.	97US-0048875P.	PR	
XX			PR 06-JUN-1997;	
CC	The invention relates to an isolated HEMAEBO protein. The protein is	97US-0048875P.	PR	
CC	useful for preparing a composition for diagnosing or treating autoimmune	PR 06-JUN-1997;		
CC	disorders e.g. multiple sclerosis and systemic lupus erythematosus;	97US-0048875P.	PR	
CC	haemopoietic cell disorders e.g. agammaglobulinaemia and ataxia	PR 06-JUN-1997;		
CC	telangiectasia; blood coagulation disorders e.g. afibrinogenemia and	97US-0048875P.	PR	
CC	thrombocytopenia; allergy; graft-versus-host disease; inflammatory	PR 06-JUN-1997;		
CC	conditions e.g. ischaemia-reperfusion injury and arthritis;	97US-0048880P.	PR	
CC	hyperproliferative disorders e.g. cancer and purpura; infectious disease	06-JUN-1997;		
CC	e.g. viral infection and bacterial infection. The polynucleotide or	97US-0048880P.	PR	
CC	protein can be used to regenerate damaged tissue e.g. ulcers and	06-JUN-1997;		
CC	Alzheimer's disease. The present sequence represents the amino acid	97US-0048883P.	PR	
CC	sequence of a novel human secreted protein. Note: The sequence data for	06-JUN-1997;		
CC	this patent did not form part of the printed specification but was	97US-0048884P.	PR	
CC	obtained in electronic format directly from USPTO at	06-JUN-1997;		
CC	http://seqdb.uspto.gov/Sequence.html?docID=20030065160	97US-0048885P.	PR	
SQ	Sequence 257 AA;	04-DEC-1998;		
	Query Match 40.3%; Score 1230; DB 6; Length 257;	98US-00205258.		
	Best Local Similarity 93.1%; Pid: No. 3.3e-110; Mismatches 3; Indels 6; Gaps 1;			
Qy	160 MAALTSHLQONQNSNNWNRLTRSKCKDVFMPSSSSELQESRGLSNFTSHILIKEDEC			
Db	1 MAALTSHLQONQNSNNWNRLTRSKCKDVFMPSSSSELQESRGLSNFTSHILIKEDEC			
	60			

PR 06-JUN-1997; 97US-0048972P.
 PR 06-JUN-1997; 97US-0048974P.
 PR 06-JUN-1997; 97US-0049019P.
 PR 06-JUN-1997; 97US-0049020P.
 PR 06-JUN-1997; 97US-0049373P.
 PR 06-JUN-1997; 97US-0049374P.
 PR 06-JUN-1997; 97US-0049375P.
 PR 05-SEP-1997; 97US-0057627P.
 PR 05-SEP-1997; 97US-0057628P.
 PR 05-SEP-1997; 97US-0057629P.
 PR 05-SEP-1997; 97US-0057634P.
 PR 05-SEP-1997; 97US-0057635P.
 PR 05-SEP-1997; 97US-0057642P.
 PR 05-SEP-1997; 97US-0057643P.
 PR 05-SEP-1997; 97US-0057644P.
 PR 05-SEP-1997; 97US-0057645P.
 PR 05-SEP-1997; 97US-0057646P.
 PR 05-SEP-1997; 97US-0057647P.
 PR 05-SEP-1997; 97US-0057648P.
 PR 05-SEP-1997; 97US-0057649P.
 PR 05-SEP-1997; 97US-0057650P.
 PR 05-SEP-1997; 97US-0057651P.
 PR 05-SEP-1997; 97US-0057654P.
 PR 05-SEP-1997; 97US-0057661P.
 PR 05-SEP-1997; 97US-0057662P.
 PR 05-SEP-1997; 97US-0057666P.
 PR 05-SEP-1997; 97US-0057667P.
 PR 05-SEP-1997; 97US-0057668P.
 PR 05-SEP-1997; 97US-0057760P.
 PR 05-SEP-1997; 97US-0057761P.
 PR 05-SEP-1997; 97US-0057762P.
 PR 05-SEP-1997; 97US-0057763P.
 PR 05-SEP-1997; 97US-0057764P.
 PR 05-SEP-1997; 97US-0057765P.
 PR 05-SEP-1997; 97US-0057770P.
 PR 05-SEP-1997; 97US-0057771P.
 PR 05-SEP-1997; 97US-0057774P.
 PR 05-SEP-1997; 97US-0057775P.
 PR 05-SEP-1997; 97US-0057776P.
 PR 05-SEP-1997; 97US-0057777P.
 PR 18-DEC-1997; 97US-0070933P.
 PR 04-JUN-1998; 98WO-US011422.
 PR 15-JUL-1998; 98US-0092931P.
 PR 30-JUL-1998; 98US-0094657P.
 PR XX
 PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 PT Young P, Greene JM, Ferrie AM, Ruben SM, Rosen CA, Hu J;
 PI Olsen HS, Ebner R, Brewer LA, Moore PA, Shi Y, Florence C;
 PI Florence K, Lafleur DW, Ni J, Pan P, Wei Y, Fischer CL, Soppet DR;
 PI Li Y, Zeng Z, Kyaw H, Yu G, Feng P, Dillon PJ, Dillon GA;
 PI Carter KC;
 XX
 WPI; 2003-511926/4B.
 DR N-PSDB; AC044672.

The invention relates to an isolated protein comprising amino acid residues 33-205 or 1-205 of a novel human secreted protein appearing as ABO2622. The protein 1 encoded by one of 238 disclosed cDNA sequences encoding 238 secreted proteins. ABO2625 is a precerebellin-like protein. Also included are a composition comprising the protein and a carrier and an isolated protein produced by expressing the protein cited above by a cell, and recovering the protein. The proteins are useful for diagnosing or treating neurodegenerative and behavioural disorders (e.g. Alzheimer's disease, Parkinson's disease, Huntington's disease, schizophrenia, mania, dementia, paranoia, psychoses or autism), immune disorders (e.g. infection, inflammation, allergy), liver disorders (e.g. hepatoblastoma, jaundice, hepatitis), immunological disorders (e.g. AIDS, leukaemia, rheumatoid arthritis, sepsis, acne, psoriasis) and cancer. The present sequence is one of the 238 disclosed novel secreted proteins. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from USPTO at: - Seqdata.uspto.gov/sequence.html?docid=6525174B1

SQ Sequence 257 AA;

Query	Match	Score	Length
Best	Local Similarity	40.3%	257
Matches	Conservative	93.1%	
Db	MATCHLONQNSNNSNWNLTRTSKCKKDVKMPSSSSESQESRGISNFTSHLLIKEDEG	219	
Db	MAALTSHLONQNSNNSNWNLTRTSKCKKDVKMPSSSSESQESRGISNFTSHLLIKEDEG	60	1
Db	VDDVNFRKVPKPGKOTILKGIPKTKKKCRKSCSGVYQDSKERSVCKADAESEPVAVD	279	61
Db	VDDVNFRKVPKPGKOTILKGIPKTKKKCRKSCSGVYQDSKERSVCKADAESEPVAVD	120	
QY	QKSQLDRTVCITSAGACGETSVTSEBBNSLUKKERSLSSGSNFCSBQTKSGIINKFCSA	339	
Db	OQSQLDRTVCISDAGGETSVTSEBBNSLUKKERSLSSGSNFCSBQTKSGIINKFCSA	180	121
QY	KDSEHNEKYEDUTPLFSEEEIGTKVVEVERKEHLHTDILKRSSEMDNNCSPTRKDFGKIF	399	
Db	KDSEHNEKYEDUTPLFSEEEIGTKVVEVERKEHLHTDILKRSSEMDNNCSPTRKDFGKIF	235	181
QY	QEDTPRTQLERKTSLYF	418	
Db	EDTPRTQLERKTSLYF	233	400
QY	EDTPRTQLERKTSLYF	233	236
QY	-EDTIERINTDRKKENPKVF	253	

RESULT 11

ID	ADD8916 standard; protein; 202 AA.
XX	ADD8916;
AC	ADD8916;
DT	29-JAN-2004 (first entry)
XX	Human 5-methylcytosine DNA glycosylase N-terminal deletion mutant.
XX	Human; 5-methylcytosine DNA glycosylase; enzyme; CpG; mutant; mutain.
XX	Homo sapiens.
OS	OS
XX	Young P, Greene JM, Ferrie AM, Ruben SM, Rosen CA, Hu J;
PN	PN WO2003078593-A2.
XX	Olsen HS, Ebner R, Brewer LA, Moore PA, Shi Y, Florence C;
PD	PD 25-SEP-2003.
XX	Florence K, Lafleur DW, Ni J, Pan P, Wei Y, Fischer CL, Soppet DR;
PF	PF 14-MAR-2003; 2003WO-US007933.
XX	Carter KC;
PR	PR 15-MAR-2002; 2002US-0364689P.
XX	(EPIC-) ERGENOMICS AG.
PA	PA Lofton-Day CE, Day JK;
XX	DR WPI; 2003-779127/73.

PT labeling methylated or methylatable CpG sequences, useful e.g. for diagnostic detection of altered methylation, comprises replacing methylated cytosine by labeled cytosine.

PT Claim 11; Page 71-72; 73pp; English.

CC The present sequence is the protein sequence of an N-terminal deletion mutant of human 5-methylcytosine DNA glycosylase (5-MCnG), in which amino

acid residue 1 corresponds to amino acid 379 of the full-length protein ADD9906. 5-McDG acts by cleaving glycosylic bonds at methylated CpG sites of DNA, removing 5-methylcytosine from the DNA backbone as a free base. The N-terminal deletion mutant shows enhanced deglycosylase specificity towards Cpg dinucleotide sequences. Human 5-McDG can be used in a claimed method for labelling Cpg sequences corresponding to methylated Cpg sequences in an isolated DNA sample. The method comprises: digesting the genomic DNA with a restriction endonuclease to produce genomic DNA fragments; treating the genomic DNA fragments with 5-McDG such that one or more 5-methylcytosine bases are removed to produce abasic genomic DNA fragments; and treating these abasic genomic fragments with base excision repair enzymes in the presence of labelled dCtp such that 5-methylcytosine removed from the genomic DNA fragments by 5-McDG is replaced by labelled cytosine in the one or more corresponding positions of the abasic genomic DNA fragments to produce labelled genomic DNA fragments, so that specific labelling of Cpg sequences corresponding to methylated Cpg sequences is achieved. The 5-McDG is also used in a claimed method for comparing Cpg methylation status, extent or pattern between or among reference and test genomic DNA samples, and in a claimed method for labelling potentially-methylatable Cpg sequences in Cpg-containing genomic DNA, including comparison of methylation pattern between healthy and diseased samples, for diagnosis.

Sequence 202 AA;

Query Match 36.2%; Score 1106; DB 7; Length 202;
Best Local Similarity 100.0%; Pred. No. 2.4e-98; Mismatches 0; Indels 0; Gaps 0;

Db 1 GSEMDDNCSPTRKDFGEKIFQEDTIPRQERRKTSLYFSKYNKEALSPPRKAKWW 438
Qy 379 GSEMDDNCSPTRKDFGEKIFQEDTIPRQERRKTSLYFSKYNKEALSPPRKAKWW 438
Db 61 TPPRSPEPNLVQETLFHDPKULLATIFLNRTSKGMAIPVLUKFLEKYPASAVARTADWRD 498
Qy 499 VSELURKPLGIYDLRAKTIIVKESDEYLTKOMKPYIHLHGIGKYGNDSYRIFCVNEMKQHP 558
Db 121 VSELURKPLGIYDLRAKTIIVKESDEYLTKOMKPYIHLHGIGKYGNDSYRIFCVNEMKQHP 180
Qy 559 EDHKUNKYHDLWENHEKLIS 580
Db 181 EDHKUNKYHDLWENHEKLIS 202

Query Match 28.8%; Score 880.5; DB 7; Length 416;
Best Local Similarity 78.2%; Pred. No. 6.3e-76; Mismatches 21; Indels 4; Gaps 2;

Db 210 RDSADGVDWSPSDKSFKTFAVQARPTESAPRQVDRKTSYFSSKYSKEALSPPRK 269
Qy 434 AFKKWTTPRSPEPNLVQETLFHDPKULLATIFLNRTSKGMAIPVLUKFLEKYPASAVRT 493
Db 270 AFRKWTTPRSPEPNLVQETLFHDPKULLATIFLNRTSKGMAIPVLUKFLEKYPASAVRT 329

Qy 494 ADWRDVSLLKPLGIYDLRAKTIIVKFSDEYLTKOMKPYIHLHGIGKYGNDSYRIFCVNEM 553
Db 330 ADWKMSLLKPLGIYDLRAKTIIVKFSDEYLTKOMKPYIHLHGIGKYGNDSYRIFCVNEM 389
Qy 554 KQVRPDKUNKYHDLWENHEKLISL 579
Db 390 KEVOPQDHKUNYHDLWENHEKLISV 415

RESULT 12
ADD9908
ID ADD9908 standard; protein; 416 AA.

AC ADD9908;
XX DT 29-JAN-2004 (first entry)
XX Chicken 5-methylcytosine DNA glycosylase.
XX
XX Chicken; 5-methylcytosine DNA glycosylase; enzyme; Cpg.
XX OS Gallus gallus.
XX PN WO2003/078593-A2.
XX PD 25-SEP-2003.
XX
XX 14-MAR-2003; 2003WO-US007933.
XX PR 15-MAR-2002; 2002US-0364689P.
XX PA (EPIC-) EPIGENOMICS AG.
XX
XX PI Lofton-Day CB, Day JK;

XX WPI; 2003-779127/73.
DR N-ISDB; ADD9907.

XX PT Labeling methylated or methylatable Cpg sequences, useful e.g. for diagnostic detection of altered methylation, comprises replacing PT methylated cytosine by labeled cytosine.

XX PS Claim 11; Page 56-58; 73pp; English.

The present sequence is the protein sequence of chicken 5-methylcytosine DNA glycosylase (5-McDG). The enzyme acts by cleaving glycosylic bonds at methylated Cpg sites of DNA, removing 5-methylcytosine from the DNA backbone as a free base. Chicken 5-McDG can be used in a claimed method for labelling Cpg sequences corresponding to methylated Cpg sequences in an isolated DNA sample. The method comprises: digesting the genomic DNA with a restriction endonuclease to produce genomic DNA fragments; treating the genomic DNA fragments with 5-McDG such that one or more 5-methylcytosine bases are removed to produce abasic genomic DNA fragments; and treating these abasic genomic DNA fragments by base excision repair enzymes in the presence of labelled dCtp such that 5-methylcytosine removed from the genomic DNA fragments by 5-McDG is replaced by labelled cytosine in the one or more corresponding positions of the abasic genomic DNA fragments to produce labelled genomic DNA fragments, so that specific labelling of Cpg sequences corresponding to methylated Cpg sequences is achieved. The 5-McDG is also used in a claimed method for comparing Cpg methylation status, extent or pattern between or among reference and test genomic DNA samples, and in a claimed method for labelling potentially-methylatable Cpg sequences in Cpg-containing genomic DNA fragments. The methods are used to identify methylated and/or potentially methylatable Cpg dinucleotides in genomic DNA, including comparison of methylation pattern between healthy and diseased samples, for diagnosis.

Sequence 416 AA;

Query Match 28.8%; Score 880.5; DB 7; Length 416;
Best Local Similarity 78.2%; Pred. No. 6.3e-76; Mismatches 21; Indels 4; Gaps 2;

Db 210 RDSADGVDWSPSDKSFKTFAVQARPTESAPRQVDRKTSYFSSKYSKEALSPPRK 269
Qy 434 AFKKWTTPRSPEPNLVQETLFHDPKULLATIFLNRTSKGMAIPVLUKFLEKYPASAVRT 493
Db 270 AFRKWTTPRSPEPNLVQETLFHDPKULLATIFLNRTSKGMAIPVLUKFLEKYPASAVRT 329

Qy 494 ADWRDVSLLKPLGIYDLRAKTIIVKFSDEYLTKOMKPYIHLHGIGKYGNDSYRIFCVNEM 553
Db 330 ADWKMSLLKPLGIYDLRAKTIIVKFSDEYLTKOMKPYIHLHGIGKYGNDSYRIFCVNEM 389
Qy 554 KQVRPDKUNKYHDLWENHEKLISL 579
Db 390 KEVOPQDHKUNYHDLWENHEKLISV 415

RESULT 13
ADD9917
ID ADD9917 standard; protein; 147 AA.

AC ADD9917;
XX DT 29-JAN-2004 (first entry)
XX Human 5-methylcytosine DNA glycosylase N-terminal deletion mutant.
XX KW Human; 5-methylcytosine DNA glycosylase; enzyme; Cpg; mutant; mutein.
XX OS Homo sapiens.
XX PN WO2003/078593-A2.
XX PD 25-SEP-2003.

XX
PP 14-MAR-2003; 2003WO-US007933.
XX
PR 15-MAR-2002; 2002US-0364689P.
XX
PA (EPIC-) EPIGENOMICS AG.
XX
PI Lofton-Day CB, Day JK;
XX
DR WPI; 2003-779127/73.

XX
PT Labeling methylated or methylatable CPG sequences, useful e.g. for diagnostic detection of altered methylation, comprises replacing PT methylated cytosine by labeled cytosine.

XX
Claim 11; Page 73; 73pp; English.

The present sequence is the protein sequence of an N-terminal deletion mutant of human 5-methylcytosine DNA glycosylase (5-MCG), in which amino acid residue 1 corresponds to amino acid 434 of the full-length protein ADD8906. 5-MCG acts by cleaving glycosidic bonds at methylated CPG sites of DNA, removing 5-methylcytosine from the DNA backbone as a free base. The N-terminal deletion mutant shows enhanced deglycosylase specificity towards CPG dinucleotide sequences. Human 5-MCG can be used in a claimed method for labelling CPG sequences corresponding to methylated CPG sequences in an isolated DNA sample. The method comprises: digesting the genomic DNA with a restriction endonuclease to produce genomic DNA fragments; treating the genomic DNA fragments with 5-MCG such that one or more 5-methylcytosine bases are removed to produce abasic genomic DNA fragments; and treating these abasic genomic DNA fragments with base excision repair enzymes in the presence of labelled dCTP such that 5-methylcytosine removed from the genomic DNA fragments by 5-MCG is replaced by labelled cytosine in the one or more corresponding positions of the abasic genomic DNA fragments to produce labelled genomic DNA fragments, so that specific labelling of CPG sequences corresponding to methylated CPG sequences is achieved. The 5-MCG is also used in a claimed method for comparing CPG methylation status, extent or pattern between or among reference and test genomic DNA samples, and in a claimed method for labelling potentially-methylatable CPG sequences in CPG-containing genomic DNA fragments. The methods are used to identify methylated and/or potentially methylatable CPG dinucleotides in genomic DNA, including comparison of methylation pattern between healthy and diseased samples, for diagnosis.

XX
Sequence 147 AA;

Query Match 26.7%; Score 816; DB 7; Length 147;
Best Local Similarity 100.0%; Pred. No. 2e-70; Length 147;
Matches 147; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 434 ATKKWTPRSPENLVLQETLFDHPWKLUJATFLNRTSGKMAIPVNLKPLEKPSAVART 493
Db 1 ARKKWTPRSPENLVLQETLFDHPWKLUJATFLNRTSGKMAIPVNLKPLEKPSAVART 60
Qy 494 AWRDVSLLKPLGLYDLRATIKVFSDEYLTKQWKPPIELAGIGKYGNDSSRIFCVNEW 553
Db 61 ADWRDVSLLKPLGLYDLRATIKVFSDEYLTKQWKPPIELAGIGKYGNDSSRIFCVNEW 120
Qy 554 KQWHPHEPKLNUCKHDWLWENHEKLISL 580
Db 121 KQWHPEDHKLNKQHDWLWENHEKLISL 147

RESULT 14

ID AAE22568 standard; peptide: 68 AA.

AC AAE22568;

DT 26-JUL-2002 (first entry)

XX
DE Human MB4 methyl CPG binding motif.

KW Gene expression; cellular chromatin; methyl CPG binding domain; cancer; localisation domain; diabetic retinopathy; ischaemia; HIV infection; KW human immuno deficiency virus; macular degeneration; vascular disease; KW rheumatoid arthritis; psoriasis; Alzheimer's disease; muscular dystrophy; KW sickle cell anaemia; stroke; neurodegenerative disease; cratic fibrosis; KW gene therapy; cytoprotective; antidiabetic; vasotropic; KW neuroprotective; nootropic; cerebroprotective; antibacterial; antifungal; KW antiviral; human; MB4 methyl CPG binding motif.

Homo sapiens.

XX	Key	Location/Qualifiers
XX	Region	1..3 /label= Beta1_helix
XX	Region	10..15 /label= Beta2_helix
XX	Region	16..26 /label= L1_loop
XX	Region	27..32 /label= Beta3_helix
XX	Region	36..38 /label= Beta4_helix
XX	Region	42..48 /label= Alpha1_helix
XX	Region	49..60 /label= L2_ loop
XX	Region	61..68 /label= Hairpin_loop

PN WO200226950-A2.

XX
PD 04-APR-2002.
XX
PP 28-SEP-2001; 2001WO-US042377.
XX
PR 29-SEP-2000; 2000US-023884P.
XX
(SANG-) SANGAMO BIOSCIENCES INC.
XX
Wolffe AP, Urnov F, Lai A, Raschke B;
XX
DR WPI; 2002-372124/40.

XX
PT Compartmentalizing a region of interest in cellular chromatin which facilitates the modulation of the expression of a gene comprises contacting the gene with a composition comprising a localization domain and a DNA binding domain.

XX
PS Example 2; Fig 1A; 85pp; English.

The present invention relates to methods and compositions for regulating gene expression. In particular the method of compartmentalising a region of interest in cellular chromatin comprises contacting the region of interest with a composition that binds to a binding site in cellular chromatin, where the binding site is in a gene of interest and the composition comprises a localization domain (e.g., methyl CPG binding domain obtained from MeCP2, MBP1, MBP2, MBP3, dmBDP-like domain and a DNA binding domain (or functional fragment)). The method is useful for compartmentalising a region of interest in cellular chromatin which facilitates the modulation of the expression of a gene using a fusion molecule comprising a DNA binding domain and a localization domain that binds to the chromatin. The fusion molecules or polypeptides can be used to prepare pharmaceutical compositions to prevent or treat cancer, ischaemia, diabetic retinopathy, macular degeneration, HIV infection, rheumatoid arthritis, psoriasis, sickle cell anaemia, vascular disease, Alzheimer's disease, muscular dystrophy, neurodegenerative diseases, cystic fibrosis, stroke, bacterial, viral or fungal infections. Sequences of the invention are also used in gene therapy. The present sequence is exemplification of the invention

XX
SQ sequence 68 AA;

Query Match 11.7%; Score 357; DB 5; Length 68;
 Best Local Similarity 98.5%; Pred. No. 2.1e-26; Matches 67; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 81 ECRKSVPCGMRVQRQLFCKTAGRFDVFISPOGLKFRSKSSLANTLHKNGETSLKPED 140
 1 ECRKSVPCGMRVQRQLFCKTAGRFDVFISPOGLKFRSKSSLANTLHKNGETSLKPED 60

OY 141 FDFTVLSK 148
 61 FDFTVLSK 68

DB

RESULT 15

ABBS1147

ID ABBS1147 standard; protein; 50 AA.

XX

AC ABBS1147;

XX DT 07-FEB-2002 (first entry)

XX DE Human secreted protein encoded by gene 168 SEQ ID NO:1100.

XX KW Human; secreted protein; immunomodulatory; antisclerotic; anti-HIV; dermatological; immunosuppressive; antiinflammatory; immunostimulant; cyclostatic; cardiant; vascular; anti-angiogenic; ophthalmological; neuroprotective; nootropic; anticonvulsant; anti-alzheimers; vulnerary; anti-parkinsonian; antimicrobial; gene therapy; vaccine; immune disorder; multiple sclerosis; systemic lupus erythematosus; HIV infection; cancer; human immunodeficiency virus; hyperproliferative disorder; wound healing; Gaucher's disease; cardiovascular disease; Scimitar syndrome; chemotaxis; Chaga's cardiomyopathy; coronary arteriosclerosis; angiogenic disorder; cornel graft neovascularisation; diabetic retinopathy; angiogenic disorder; neurological disorder; Huntington's chorea; Alzheimer's disease; Parkinson's disease; infectious disease; chromosome 3.

XX OS Homo sapiens.

XX PN WO200162891-A2.

XX PD 30-AUG-2001.

XX PR 21-FEB-2001; 2001WO-US005614.

XX PR 24-FEB-2000; 2000US-0184836P.

XX PR 29-MAR-2000; 2000US-0193170P.

XX PA (HUMA-) HUMAN GENOME SCI INC.

XX Ni J, Elbner R, Lafleur DW, Moore PA, Olsen HS, Rosen CA,
 PI Ruben SM, Soppet DR, Young PE, Shi Y, Florence KA, Wei Y,
 PI Florence C, Hu J, Li Y, Kyaw H, Fischer CL, Ferrie AM, Fan P,
 PI Feng P, Endress GA, Dillon PJ, Carter KC, Brewer LA, Yu G, Zeng Z,
 PI Greene JM;
 XX DR MPI; 2001-625724/72.

XX PT Nucleic acids encoding 207 human secreted polypeptides, useful for preventing, diagnosing and/or treating, e.g. cancers, Parkinson's disease and diabetic retinopathy.

XX PS Disclosure; Page 366; 1533PP; English.

CC ABBA0301 to ABBS1267 and ABBA3194 to ABBA8341 represent human secreted proteins (I) and polyneurotide (II) sequences. (I) and (II) have various activities based on the tissues and cells the genes are expressed in. Example of these activities include: immunomodulatory; antisclerotic; dermatological; immunosuppressive; antiinflammatory; immunostimulant; anti-HIV; cyclostatic; cardiant; anti-angiogenic; ophthalmological; neuroprotective; nootropic; anticonvulsant; anti-alzheimers; vascular; anti-parkinsonian; antimicrobial; and vulnerary. (I) and (II) can be used in gene therapy and vaccine production. (I) and (II) can be used in the prevention, diagnosis and treatment of immune disorders (e.g. multiple

Sequence 50 AA;

Search completed: August 22, 2005, 10:05:09
 Job time : 172 secs

Query Match 8.3%; Score 254; DB 4; Length 50;
 Best Local Similarity 98.0%; Pred. No. 1.3e-16; Matches 49; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 110 FISPOGLKFRSKSSLANTLHKNGETSLKPEDFDFTVLSKRGKSYKDCS 159
 1 FSSPQGLKFRSKSSLANTLHKNGETSLKPEDFDFTVLSKRGKSYKDCS 50

DB

CC sclerosis, systemic lupus erythematosus and human immunodeficiency virus (HTV) infections), hyperproliferative disorders (e.g. cancers and Gaucher's disease), cardiovascular diseases (e.g. Scimitar syndrome, Chaga's cardiomyopathy and coronary arteriosclerosis), angiogenic disorders (e.g. corneal graft neovascularisation and diabetic retinopathy), neurological disorders (e.g. Huntington's chorea, Alzheimer's disease and Parkinson's disease), infectious diseases and/or promoting wound healing, regeneration and/or chemotaxis. ABAA8185 to ABAA83193 and ABBA0300 represent sequences used in the exemplification of the present invention

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Om protein - protein search, using sw model

Run on: August 22, 2005, 10:02:09 ; Search time 163 Seconds

(without alignment(s)) 1393.374 Million cell updates/sec

US-10-629-951-2

3055

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Title: US-10-629-951-2

Perfect score:

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Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Searched: 1759131 seqs, 391586102 residues

Total number of hits satisfying chosen parameters: 1759131

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Published Applications RA: *
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	3055	100.0	580	14	US-10-389-853-2
2	3055	100.0	580	15	US-10-629-951-30
3	3011	98.6	574	15	US-10-629-951-30
4	1639	53.6	307	14	US-10-389-853-11
5	1326	43.4	384	15	US-10-629-951-11
6	1230	40.3	257	10	US-09-933-767-416
7	1230	40.3	257	14	US-10-004-860-416
8	1230	40.3	257	14	US-10-023-282-416
9	1106	36.2	202	14	US-10-389-853-12
10	880.5	28.8	416	14	US-10-389-853-4
11	816	26.7	417	14	US-10-389-853-13

ALIGNMENTS

RESULT 1 US-10-389-853-2	Query Match	100.0%; Score 3055; DB 14; Length 580;	Sequence 32, Appl
; Sequence 2, Application US/10389853	; Best Local Similarity 100.0%; Pred. No. 8.2e-236;	Sequence 30, Appl	
; Publication No. US20030180779A1	; Matches 580; Conservative 0; Mismatches 0; Indexes 0; Gaps 0;	Sequence 37, Appl	
; GENERAL INFORMATION:		Sequence 7, Appl	
; APPLICANT: Lofton-Day, Cathy E.		Sequence 5, Appl	
; APPLICANT: Day, John K.		Sequence 11, Appl	
; TITLE OF INVENTION: Discovery and Diagnostic Methods Using 5-Methylcytosine DNA Glyco		Sequence 1, Appl	
; FILE REFERENCE: 47675-36		Sequence 17, Appl	
; CURRENT APPLICATION NUMBER: US/10-389, 853		Sequence 18, Appl	
; CURRENT FILING DATE: 2003-03-14		Sequence 19, Appl	
; PRIORITY APPLICATION NUMBER: 60/364, 689		Sequence 548, Appl	
; PRIORITY FILING DATE: 2005-03-15		Sequence 550, Appl	
; NUMBER OF SEQ ID NOS: 13		Sequence 553, Appl	
; SOFTWARE: Patentin version 3.1		Sequence 8707, Appl	
; SEQ ID NO: 2		Sequence 2, Appl	
; LENGTH: 580		Sequence 7981, Appl	
; TYPE: PRT			
; ORGANISM: Homo sapiens			
US-10-389-853-2			

Query	Match	100.0%; Score 3055; DB 14; Length 580;	Sequence 32, Appl
Best Local Similarity	100.0%; Pred. No. 8.2e-236;	Sequence 30, Appl	
Matches	580; Conservative 0; Mismatches 0; Indexes 0; Gaps 0;	Sequence 37, Appl	
Sequence 2, Appl		Sequence 7, Appl	
Sequence 24, Appl		Sequence 5, Appl	
Sequence 11, Appl		Sequence 11, Appl	
Sequence 29, Appl		Sequence 1, Appl	
Sequence 416, Appl		Sequence 17, Appl	
Sequence 416, Appl		Sequence 18, Appl	
Sequence 416, Appl		Sequence 19, Appl	
Sequence 12, Appl		Sequence 548, Appl	
Sequence 4, Appl		Sequence 550, Appl	
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Sequence 13, Appl		Sequence 8707, Appl	
Sequence 13, Appl		Sequence 2, Appl	
Sequence 13, Appl		Sequence 7981, Appl	
US-10-389-853-2			

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Db 181 RSKCKDVFMPSSSELOESRGISNFSTHLLIKEDEGVDDVNRKVKPKGKVTLKG 240
Db 241 IPKTTKGCRKSGFVOSDSRKSRESVNCNKADESPEVAQSKQSLDRTVCSAGCETL 300
Qy 241 IPKTTKGCRKSGFVOSDSRKSRESVNCNKADESPEVAQSKQSLDRTVCSAGCETL 300
Db 241 IPKTTKGCRKSGFVOSDSRKSRESVNCNKADESPEVAQSKQSLDRTVCSAGCETL 300
Db 301 SVTSEBENSLVKKERSLSSGSNFNSEOKTGINKFCSAKDSENEKYDTFLESEBTG 360
Qy 301 SVTSEBENSLVKKERSLSSGSNFNSEOKTGINKFCSAKDSENEKYDTFLESEBTG 360
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Db 361 KVEVERKEHLHTDLKRGSEMDNCSPTRKDFGKEFOEDTPRTQJERKNTLYFS 420
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RESULT 2
US-10-629-951-2
; Sequence 2, Application US/10629951
; Publication No. US20040018550A1
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/10/629, 951
; CURRENT FILING DATE: 2003-07-29
; PRIOR APPLICATION NUMBER: US/09/629, 222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463, 891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053, 936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 2
; LENGTH: 580
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-10-629-951-2

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Best Local Similarity 100.0%; Pred. No. 8.2e-236; Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 3
US-10-629-951-24
; Sequence 24, Application US/10629951
; Publication No. US20040018550A1
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/10/629, 951
; CURRENT FILING DATE: 2003-07-29
; PRIOR APPLICATION NUMBER: US/09/629, 222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463, 891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053, 936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 24
; LENGTH: 574
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-10-629-951-24

Query Match 98.6%; Score 3011; DB 15; Length 574;
Best Local Similarity 99.0%; Pred. No. 2.7e-232; Matches 574; Conservative 0; Mismatches 0; Indels 6; Gaps 1; Gaps 1;
Matches 574; Conservative 0; Mismatches 0; Indels 6; Gaps 1;

RESULT 4
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; Sequence 24, Application US/10629951
; Publication No. US20040018550A1
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/10/629, 951
; CURRENT FILING DATE: 2003-07-29
; PRIOR APPLICATION NUMBER: US/09/629, 222A
; PRIOR FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463, 891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053, 936
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; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 24
; LENGTH: 574
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-10-629-951-24

Query Match 98.6%; Score 3011; DB 15; Length 574;
Best Local Similarity 99.0%; Pred. No. 2.7e-232; Matches 574; Conservative 0; Mismatches 0; Indels 6; Gaps 1; Gaps 1;
Matches 574; Conservative 0; Mismatches 0; Indels 6; Gaps 1;

Query Match 98.6%; Score 3011; DB 15; Length 574;
Best Local Similarity 99.0%; Pred. No. 2.7e-232; Matches 574; Conservative 0; Mismatches 0; Indels 6; Gaps 1; Gaps 1;
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RESULT 4
 US-10-389-853-11
 ; Sequence 11, Application US/10389853
 ; Publication No. US20040018550A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Norton-Day, John K.
 ; TITLE OF INVENTION: Discovery and Diagnostic Methods Using 5-Methylcytosine DNA Glyco-
 ; FILE REFERENCE: 47675-36
 ; CURRENT APPLICATION NUMBER: US/10/389,853
 ; CURRENT FILING DATE: 2003-03-14
 ; PRIORITY FILING DATE: 1997-03-15
 ; PRIORITY APPLICATION NUMBER: 60/364,689
 ; SEQ ID NO: 11
 ; LENGTH: 307
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 ; FEATURE: NON_TER
 ; LOCATION: (1); (1)
 ; OTHER INFORMATION: functional N-terminal deletion mutant of hMBD4 (SEQ ID NO:2); aa no. 1 corresponds to aa no. 274 of hMBD4; mutant shows enhanced d-
 ; OTHER INFORMATION: e.g.,cosylase specificity towards CpG dinucleotide sequences; see
 ; US-10-389-853-11

Query Match 53.6%; Score 1639; DB 14; length 307;
 Best Local Similarity 100.0%; Pred. No. 1e-122; Mismatches 0; Indels 0; Gaps 0;
 Matches 307; Conservative 0;

Qy 274 ESEPVAAQKSQLDRVTCTSDAGACETLSVTSEENSLVKKERSLSSNSNFCSEQKTSII 333
 Db 1 ESEPVAAQKSQLDRVTCTSDAGACETLSVTSEENSLVKKERSLSSNSNFCSEQKTSII 60
 Qy 334 NKFCSAKOSAKENEKYEDTPLESEBEGTKVEVERKEHLHTDIKRGSEMDDNCSPTRKD 393
 Db 61 NKFCSAKOSAKENEKYEDTPLESEBEGTKVEVERKEHLHTDIKRGSEMDDNCSPTRKD 120
 Qy 394 TGFKIQDQTIPRTOBERKTSLYFSKVNKEALSPPRKAFCWTPRSPPRNVLVQTLF 453
 Db 121 TGEKIQDQTIPRTOIERKTSLYFSKVNKEALSPPRKAFCWTPRSPPRNVLVQTLF 180

RESULT 5
 US-10-629-951-29
 ; Sequence 29, Application US/10629951
 ; Publication No. US20040018550A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Bellacosa, Alfonso
 ; TITLE OF INVENTION: Method for Detection of Transition
 ; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
 ; FILE REFERENCE: FCCC 96-21
 ; CURRENT APPLICATION NUMBER: US/10/629, 951
 ; CURRENT FILING DATE: 2003-07-29
 ; PRIORITY APPLICATION NUMBER: US/09/629, 222A
 ; PRIORITY FILING DATE: 2000-07-31
 ; PRIORITY APPLICATION NUMBER: 09/463, 891
 ; PRIORITY FILING DATE: 2000-01-28
 ; PRIORITY APPLICATION NUMBER: PCT/US98/15828
 ; PRIORITY FILING DATE: 1998-07-28
 ; PRIORITY APPLICATION NUMBER: 60/053, 936
 ; NUMBER OF SEQ ID NOS: 73
 ; SOFTWARE: FastSEQ for Windows Version 3.0
 ; SEQ ID NO: 29
 ; LENGTH: 384
 ; TYPE: PRT
 ; ORGANISM: Mus musculus
 ; US-10-629-951-29

Query Match 43.4%; Score 1326; DB 15; Length 384;
 Best Local Similarity 54.3%; Pred. No. 1.e-97; Mismatches 71; Indels 138; Gaps 4;
 Matches 283; Conservative 29; Mismatches 71; Indels 138; Gaps 4;

Qy 36 KEDVAMELERYGEDEQQMMIKRSSLCECNPLQPIASQAFGATAGCERKSYPCGWERVK 95
 Db 1 KEDIAVULGGVGEDEGDKLV--SSRSLLQEPST--LSSTATEGHKVPCKGWERVK 57
 Qy 96 ORLFKGKTAGRFDVYFTSPQGLKFRSKSLANYLHNGGETSLKPEDPDFTVLSKRGKRY 155
 Db 58 ORLSKGKTAGKFDVYFISPOGLKFRSKSLANYLHNGGETELKPEDPDFTVLSKRGKRY 117
 Qy 156 KDCSMLALTSHLQGNSNNNLRTSKCKDVFENPSSSELOSSRGLNFNSTHLLK 215
 Db 118 KHOSLAALTSQPNETVSKONLKURSKWKDVLPLSGTSESPSSGLSNNSACILLR 177
 Qy 216 EDEGVDDVNFRKVKKGKVTLKG1FPIKKKKGCRKCSGCFVQDSKRSVCNKADES 275
 Db 178 EHDQDVSERKRRKVKVTLKGASOKRKSLBOSTNRKRAS----- 228
 Qy 276 ESEPVAAQKSQLDRVTCTSDAGACETLSVTSEENSLVKKERSLSSNSNFCSEQKTSII 335
 Db 229 ----- 228

Qy 336 FCSAKOSAKENEKYEDTPLESEBEGTKVEVERKEHLHTDIKRGSEMDDNCSPTRKD 395
 Db 229 ----- 228

QY 456 PWKLLIATIINRISGRMAIEPVVLUKFLEKPSAETWADWRDVSELLAKPLGLYDLRAKT 515
 PRIOR APPLICATION NUMBER: 60/048, 916
 PRIOR FILING DATE: 1997-06-05
 Db 284 PWKLLIATIIFARTSKMAMIPVLMFLEKPSAETWADWRDVSELLUKLGLYDLRAKT 343
 PRIOR APPLICATION NUMBER: 60/049, 373
 PRIOR FILING DATE: 1997-06-05
 QY 516 IVKFSDEYLTYLKWYKTYIELIGIG-KYGNDSYRIFCVNEWHQ 555
 PRIOR APPLICATION NUMBER: 60/048, 875
 PRIOR FILING DATE: 1997-06-05
 Db 344 1IKFSDEYLTKWYKTYIELIGIG-KYGNDSYRIFCVNEWHQ 384
 PRIOR APPLICATION NUMBER: 60/049, 374
 PRIOR FILING DATE: 1997-06-05
 RESULT 6 US-09-933-767-416
 Sequence 416 Application US/09933767
 Publication No. US20030181692A1
 GENERAL INFORMATION:
 APPLICANT: Ni et al.
 TITLE OF INVENTION: 207 Human Secreted Proteins
 FILE REFERENCE: P2007P2
 CURRENT APPLICATION NUMBER: US/09/933, 767
 CURRENT FILING DATE: 2001-08-22
 PRIOR APPLICATION NUMBER: PCT/US01/05614
 PRIOR FILING DATE: 2001-02-21
 PRIOR APPLICATION NUMBER: 60/184, 836
 PRIOR FILING DATE: 2000-02-24
 PRIOR APPLICATION NUMBER: 60/193, 170
 PRIOR FILING DATE: 2000-03-29
 PRIOR APPLICATION NUMBER: 09/205, 258
 PRIOR FILING DATE: 1998-12-04
 PRIOR APPLICATION NUMBER: PCT/US98/11422
 PRIOR FILING DATE: 1998-06-04
 PRIOR APPLICATION NUMBER: 60/048, 885
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/049, 375
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/049, 881
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 880
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 896
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/049, 020
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 881
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 876
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 895
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 884
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 876
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 894
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 971
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 964
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 882
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 899
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 893
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 900
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 901
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 892
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 915
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/049, 019
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 970
 PRIOR FILING DATE: 1997-06-06
 PRIOR APPLICATION NUMBER: 60/048, 972
 ; PRIOR FILING DATE: 1997-06-06
 ; SEQ ID NO 416
 LENGTH: 257
 TYPE: PRT
 ORGANISM: Homo sapiens
 FEATURE:
 NAME/KEY: SITE
 LOCATION: (100)
 OTHER INFORMATION: xaa equals any of the naturally occurring L-amino acids
 US-09-933-767-416
 Query Match Score 40.3%; DB 10; Length 257;
 Best Local Similarity 93.1%; Pred. No. 4 6e-90; Mismatches 9; Indels 6; Gaps 1;
 Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;
 QY 160 MAALTSHIQNQSNNSNWTRSKCKKUVFMPSSSSBLQESRGLSNFTSHULLKEBEG 219
 ;

APPLICANT: Young et al.; TITLE OF INVENTION: 1997 Human Secreted Proteins FILE REFERENCE: P2007P1

CURRENT APPLICATION NUMBER: US/11/023, 282 CURRENT FILING DATE: 2011-12-20 EARLIER APPLICATION NUMBER: 09/205, 258 EARLIER FILING DATE: 1998-12-04 EARLIER APPLICATION NUMBER: PCT/US98/11422 EARLIER FILING DATE: 1998-06-04 EARLIER APPLICATION NUMBER: 60/048, 885 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/049, 375 EARLIER FILING DATE: 1997-06-05 EARLIER APPLICATION NUMBER: 60/048, 881 EARLIER FILING DATE: 1997-06-05 EARLIER APPLICATION NUMBER: 60/048, 880 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 896 EARLIER FILING DATE: 1997-06-05 EARLIER APPLICATION NUMBER: 60/049, 020 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 876 EARLIER FILING DATE: 1997-06-05 EARLIER APPLICATION NUMBER: 60/048, 895 EARLIER FILING DATE: 1997-06-05 EARLIER APPLICATION NUMBER: 60/048, 884 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 894 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 971 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 964 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 882 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 899 EARLIER FILING DATE: 1997-06-05 EARLIER APPLICATION NUMBER: 60/048, 893 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 900 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 901 EARLIER FILING DATE: 1997-06-05 EARLIER APPLICATION NUMBER: 60/048, 892 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 915 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/049, 019 EARLIER FILING DATE: 1997-06-05 EARLIER APPLICATION NUMBER: 60/048, 970 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 972 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 916 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 973 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 875 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/049, 374 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 917 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 893 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 897 EARLIER FILING DATE: 1997-06-06 EARLIER APPLICATION NUMBER: 60/048, 898

RESULT 7
US-10-004,860-416
; Sequence 416, Application US/10004860
; Publication No. US20030065160A1
; GENERAL INFORMATION:
; APPLICANT: Young et al.
; TITLE OF INVENTION: 207 Human Secreted Proteins
; FILE REFERENCE: P2007P1
; CURRENT APPLICATION NUMBER: US/10/004,860
; CURRENT FILING DATE: 2001-12-07
; PRIORITY CLAIMS:
; Prior Application removed - See File Wrapper or Palm
NUMBER OF SEQ ID NOS: 1227
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO: 416
LENGTH: 257
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: SITE
LOCATION: (100)
OTHER INFORMATION: xaa equals any of the naturally occurring L-amino acids
US-10-004-860-416
Query Match 40.3% Score 1230; DB 14; Length 257;
Best Local Similarity 93.1%; Pred. No. 4.6e-90; 9; Indels 6; Gaps 1;
Matches 241; Conservative 3; Mismatches 1;
QY 160 MAALTHLQNSNNWURTRSKKDVFMPSSSELOESRGLSNFTSTHLLIKEDEG 219
Db 1 MAALTHLQNSNNWURTRSKKDVFMPSSSELOESRGLSNFTSTHLLIKEDEG 60
QY 220 VDDVNRKVRKPKPGKVTILKGIPRKTKKGCRKSGSGFWQDSRSKEVSNCNADESEPA 279
Db 61 VDDVNRKVRKPKPGKVTILKGIPRKTKKGCRKSGSGFWQDSRSKEVSNCNADESEPA 120
QY 280 QKSQLDRTWCISDAGAGETLSVSEENSLVKCKERSLSSGSNCFSEOKTSGINKFCSA 339
Db 121 QKSQLDRTWCISDAGAGETLSVSEENSLVKCKERSLSSGSNCFSEOKTSGINKFCSA 180
QY 340 KDSERNEKEDTFLSEEIGTKVVERKEHLHTDLKGSEMNDNCSPTRKDFGEKIP 399
Db 181 KDSERNEKEDTFLSEEIGTKVVERKEHLHTDLKGSEMNDNCSPTRKDFGEKIP 235
QY 400 QEDTPRQTERRKTSLYF 418
Db 236 -EDTPRNTDRKKENKPVF 253

RESULT 8
US-10-023-282-416
; Sequence 416, Application US/10023282
; Publication No. US20030092893A1
; GENERAL INFORMATION:

; OTHER INFORMATION: functional N-terminal deletion mutant of hMBD4 (SEQ ID NO:2); aa
; OTHER INFORMATION: no. 1 corresponds to aa no. 379 of hMBD4 protein; mutant shows en
; OTHER INFORMATION: hanced glycosylase specificity towards CpG dinucleotide sequenc
; OTHER INFORMATION: es; see Zhu et al. Nuc. Acid Res. 28:4157-4165, 2000.
; US-10-389-853-12

EARLIER APPLICATION NUMBER: 60/048, 962
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048, 963
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048, 877
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048, 878
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/070, 923
; EARLIER FILING DATE: 1997-12-18
; EARLIER APPLICATION NUMBER: 60/092, 921
; EARLIER FILING DATE: 1998-07-15
; EARLIER APPLICATION NUMBER: 60/094, 657
; EARLIER FILING DATE: 1998-07-30
; NUMBER OF SEQ ID NOS: 1227
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO: 416
; LENGTH: 257
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; LOCATION: (100)
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids

US-10-023-282-416

Query Match 40.3%; Score 1230; DB 14; Length 257;
; Best Local Similarity 93.1%; Pred. No. 4.6e-90;
; Matches 241; Conservative 3; Mismatches 9; Indels 6; Gaps 1;

Db 160 MAALTSHLQOQSNSNWNLTTRSKCKDVFMPRSSSELOEQRGLSNFSTHILKEDEG 219
; 1 MAALTSHLQLONGNSNNWNLTTRSKCKDVFMPSSSELOEQRGLSNFTTHILKEDEG 60

Qy 220 VDDVAVRKVRPKPGKVTILKGKIPKTKGGKRSKSSGFVQDSKRSKESVNKADESEPYA 279
; 61 VDDVAVRKVRPKPGKVTILKGKIPKTKGGKRSKSSGFVQDSKRSKESVNKADESEPYA 120

Db 280 QKSQDRTVCSIDAGGETTSVTSBENSLYKKERSLSQSGSNFCSEOKTGTTGINKFCSA 339
; 121 QKSQDRTVCSIDAGGETTSVTSBENSLYKKERSLSQSGSNFCSEOKTGTTGINKFCSA 180

Qy 340 KDSERNEKYEDTFLESEEIGKVVERKELHTDILKGSSEMNDNCSPTRKDPIGEKIF 399
; 181 KDSERNEKYEDTFLESEEIGKVVERKELHTDILKGSSEMNDNCSPTRKDFT---- 235

Qy 400 QDTIPTQIERRKTSLYF 418
; Db 236 -EDTIPRNTDRKKENPKPV 253

RESULT 9
; Sequence 12, Application US/10389853
; Publication No. US20030180779A1
; GENERAL INFORMATION:
; APPLICANT: Lofton-Day, Cathy E.

; TITLE OF INVENTION: Discovery and Diagnostic Methods Using 5-Methylcytosine DNA Glyco-

; CURRENT APPLICATION NUMBER: US/10/389, 853
; PRIORITY FILING DATE: 2003-03-14
; PRIOR APPLICATION NUMBER: 60/364, 689
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO: 4
; LENGTH: 416
; TYPE: PRT
; ORGANISM: Gallus gallus

US-10-389-853-4

Query Match 28.8%; Score 880.5; DB 14; Length 416;
; Best Local Similarity 78.2%; Pred. No. 9.6e-62;
; Matches 161; Conservative 20; Mismatches 21; Indels 4; Gaps 2;

Db 378 RGSEMDNNCS--PDKDFTGEGKIQF-EDTIPRQIERRKTSLYFSSKCNKEALSPPRK 433
; 210 RDSAADGVSWPSDPDKSFTAVOPRGTESAPTOVDRKTSYFSSKCNKEALSPPRK 269

Qy 434 AFKWTTPRSPSPNIVQETLFHDWKLLATIFARTSGKMAIPVWLKELEKPSAERT 493
; Db 270 AFRKWTTPRSPSPNIVQETLFHDWKLLATIFARTSGKMAIPVWLKELEKPSAERT 329

Qy 494 ADWDSSEILKPGIYDRAKTIKVSFBSYLTQKVKIELHGIGKCGNDSYIFCVNEW 553
; Db 330 ADWKEMSELRLRPLGIGYIYRAKTIKFSDBYLNKQVKIELHGIGKCGNDSYIFCVNEW 389

Qy 554 KQVHPEDHKLKYDHMLMENHEKLS 579
; Db 390 KEVQPODHKLNYIHWLWENHEKLSV 415

RESULT 11
; Sequence 13, Application US/10389853
; Publication No. US20030180779A1

; OTHER INFORMATION: functional N-terminal deletion mutant of hMBD4 (SEQ ID NO:2); aa
; OTHER INFORMATION: no. 1 corresponds to aa no. 379 of hMBD4 protein; mutant shows en
; OTHER INFORMATION: hanced glycosylase specificity towards CpG dinucleotide sequenc
; OTHER INFORMATION: es; see Zhu et al. Nuc. Acid Res. 28:4157-4165, 2000.
; US-10-389-853-13

ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: NON_TER
; LOCATION: (1) .. (1)

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; GENERAL INFORMATION:
; APPLICANT: Lofton-Day, Cathy E.
; APPLICANT: Day, John K.
; TITLE OF INVENTION: Discovery and Diagnostic Methods Using 5-Methylcytosine DNA Glyc
; FILE REFERENCE: 47675_36
; CURRENT FILING DATE: 2003-03-14
; PRIOR APPLICATION NUMBER: 60/364, 689
; PRIOR FILING DATE: 2002-03-15
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: PatentIn version 3.1
SEQ ID NO 13
LENGTH: 147
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE: NON_TER
NAME/KEY: NON_TER
LOCATION: (1)..(1)
OTHER INFORMATION: functional N-terminal deletion mutant of hMBD4 (SEQ ID No:2); aa OTHER INFORMATION: no. 1 corresponds to aa no. 434 of hMBD4 protein; mutant shows enhanced deglycosylase specificity towards CpG dinucleotide sequence OTHER INFORMATION: es; see Zhu et al. Nuc. Acid Res. 28:4157-4165, 2000.
US-10-389-853-13

Query Match 26.7%; Score 816; DB 14; Length 147;
Best Local Similarity 100.0%; Pred. No. 3.1e-57; Matches 147; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Db 1 AFKKWTPPRSPNLUQETLFPWKLIATPLNRTSGKMAIPVNLKFLRCPYSAEART 493
Qy 434 APKKWTPPRSPNLUQETLFPWKLIATPLNRTSGKMAIPVNLKFLRCPYSAEART 493
Db 1 AFKKWTPPRSPNLUQETLFPWKLIATPLNRTSGKMAIPVNLKFLRCPYSAEART 60
Qy 494 ADWRDVSELLKLQLGLYDRLAKTIVKVSDEYLTKQWKPPIELHGIGKYGNDSRIFCUNEW 553
Db 61 ADWRDVSELLKLQLGLYDRLAKTIVKVSDEYLTKQWKPPIELHGIGKYGNDSRIFCUNEW 120
Qy 554 KQVHPEDHKLNKQHDWLWENHEKLSSL 580
Db 121 KQVHPEDHKLNKQHDWLWENHEKLSSL 147

RESULT 12
US-10-629-951-32
Sequence 32, Application US/10629951
Publication No. US20040018550A1
GENERAL INFORMATION:
APPLICANT: Bellacosa, Alfonso
TITLE OF INVENTION: Methods for Detection of Transition
FILE REFERENCE: FCCC 96-21
CURRENT APPLICATION NUMBER: US/10/629, 951
TITLE OF INVENTION: Methods for Detection of Transition
FILE REFERENCE: FCCC 96-21
CURRENT APPLICATION NUMBER: US/10/629, 951
CURRENT FILING DATE: 2003-07-29
PRIOR APPLICATION NUMBER: US/09/629, 222A
PRIOR FILING DATE: 2000-07-31
PRIOR APPLICATION NUMBER: 09/463, 891
PRIOR FILING DATE: 2000-01-28
PRIOR APPLICATION NUMBER: PCT/US98/15828
PRIOR APPLICATION NUMBER: 60/053, 936
PRIOR FILING DATE: 1998-07-28
NUMBER OF SEQ ID NOS: 73
SOFTWARE: FastSEQ for Windows Version 3.0
SEQ ID NO 30
LENGTH: 119
TYPE: PRT
ORGANISM: Homo sapiens
US-10-629-951-30

Query Match 20.2%; Score 616; DB 15; Length 119;
Best Local Similarity 100.0%; Pred. No. 2.5e-41; Matches 119; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Db 36 KEDVAMELERYGDEBQOMMIRKSSCENPQIOPIASAQFGATAGTECKRSVPGHMRVK 95
Qy 36 KEDVAMELERYGDEBQOMMIRKSSCENPQIOPIASAQFGATAGTECKRSVPGHMRVK 95
Db 1 KEDVAMELERYGDEBQOMMIRKSSCENPQIOPIASAQFGATAGTECKRSVPGHMRVK 60
Qy 96 QRLFGKTAGRDVYFISPOQIKFRKSLSANYLHNGETSLKPELPFDFTVLKGKISR 154
Db 61 QRLFGKTAGRDVYFISPOQIKFRSKSLANYLHNGETSLKPELPFDFTVLKGKISR 119

RESULT 14
US-10-629-951-37
Sequence 37, Application US/10629951
Publication No. US20040018550A1
GENERAL INFORMATION:
APPLICANT: Bellacosa, Alfonso
TITLE OF INVENTION: Methods for Detection of Transition
FILE REFERENCE: FCCC 96-21
CURRENT APPLICATION NUMBER: US/10/629, 951
CURRENT FILING DATE: 2003-07-29
PRIOR APPLICATION NUMBER: US/09/629, 222A
PRIOR FILING DATE: 2000-07-31
PRIOR APPLICATION NUMBER: 09/463, 891
PRIOR FILING DATE: 2000-01-28
PRIOR APPLICATION NUMBER: PCT/US98/15828
PRIOR APPLICATION NUMBER: 60/053, 936
PRIOR FILING DATE: 1998-07-28
SOFTWARE: FastSEQ for Windows Version 3.0
SEQ ID NO 32
LENGTH: 126
TYPE: PRT
ORGANISM: Homo sapiens
US-10-629-951-32

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Query Match 22.8%; Score 697; DB 15; Length 126;
Best Local Similarity 100.0%; Pred. No. 8.6e-48; Matches 126; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 455 DPWKLLIATIFLNRSGKMAIPVNLKFLRCPYSAEARTADWRDVSELLKLQLGLYDLRAK 514

```

```

; SEQ ID NO: 37
; LENGTH: 85
; ORGANISM: Homo sapiens
US-10-629-951-37

Query Match          14.5%; Score 443; DB 15; Length 85;
Best Local Similarity 100.0%; Pred. No. 1.1e-27;
Matches      85; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy       69 RASAQGATAGTECRKSVPCCWERYVKQRQFLGKTAGRFDYFTISPOGLKRRSKSSLANYL 128
Db       1 RASAQGATAGTECRKSVPCCWERYVKQRQFLGKTAGRFDYFTISPOGLKFRSKSSLANYL 60
Qy       129 HKNGETSLKPBDFFTVLSKGIKS 153
Db       61 HKNGETSLKPBDFFTVLSKGIKS 85

RESULT 15
US-09-967-869A-7
; Sequence 7, Application US/09967869A
; Publication No. US20030082552A1
; GENERAL INFORMATION:
; APPLICANT: WOLFFE, Alan P.
; APPLICANT: URNOV, Fyodor
; APPLICANT: LAI, Albert
; APPLICANT: RASCHKE, Eva
; TITLE OF INVENTION: MODULATION OF GENE EXPRESSION USING LOCALIZATION
; DOMAIN: DOMAINS
; FILE REFERENCE: 8325-0019 / S19
; CURRENT APPLICATION NUMBER: US/09/967, 869A
; CURRENT FILING DATE: 2001-09-28
; NUMBER OF SEQ ID NOS: 17
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO: 7
; LENGTH: 68
; TYPE: PRT
; FEATURE:
; ORGANISM: Artificial Sequence
; OTHER INFORMATION: Description of Artificial Sequence: MBD4
US-09-967-869A-7

Query Match          11.7%; Score 357; DB 10; Length 68;
Best Local Similarity 98.5%; Pred. No. 6.2e-21;
Matches      67; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy       81 BCKRSVPCCWERYVKQRQFLGKTAGRFDYFTISPOGLKFRSKSSLANYLHKNGETSLKPED 140
Db       1 BCKRSVPCCWERYVKQRQFLGKTAGRFDYFTISPOGLKFRSKSSLANYLHKNGETSLKPED 60
Qy       141 PDFTVLSK 148
Db       61 PDFTVLSK 68

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Search completed: August 22, 2005, 10:12:39
 Job time : 164 secs

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On protein - protein search, using bw model

Run on: August 22, 2005, 10:02:09 ; Search time 43 seconds
 (without alignments)

Scoring table: BLASTm62
 GapOp 10.0 , GapExt 0.5

Searched: 513545 Beq8, 74649064 residues

Total number of hits satisfying chosen parameters: 513545

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
 Maximum Match 100%
 Listing first 45 summaries

Database :

- 1: Issued_Patents_AA;*
- 2: /cgmn_6/pctodata/1/iaa/5A_COMB_pep;*
- 3: /cgmn_6/pctodata/1/iaa/6A_COMB_pep;*
- 4: /cgmn_6/pctodata/1/iaa/6B_COMB_pep;*
- 5: /cgmn_6/pctodata/1/iaa/PCTUS_COMB_pep;*
- 6: /cgmn_6/pctodata/1/iaa/backfile1.pep;*

Pred. No. 18 is the number of result predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match Length	DB ID	Description
1	3055	100.0	580	Sequence 38, Appl
2	3055	100.0	580	Sequence 2, Appl
3	3055	100.0	580	Sequence 56, Appl
4	3011	98.6	574	Sequence 24, Appl
5	1326	43.4	384	Sequence 29, Appl
6	1230	40.3	257	Sequence 416, Appl
7	697	22.8	126	Sequence 32, Appl
8	616	20.2	119	Sequence 30, Appl
9	443	14.5	9	Sequence 37, Appl
10	254	8.3	50	Sequence 110, Ap
11	220.5	7.2	467	Sequence 69, Ap
12	220.5	7.2	467	Sequence 70, Ap
13	213.5	7.0	486	Sequence 74, Appl
14	209	6.8	476	Sequence 75, Appl
15	208.5	6.8	477	Sequence 63, Appl
16	208.5	6.8	486	Sequence 55, Appl
17	208.5	6.8	486	Sequence 59, Appl
18	208.5	6.8	486	Sequence 60, Appl
19	208.5	6.8	486	Sequence 62, Appl
20	208.5	6.8	486	Sequence 65, Appl
21	208.5	6.8	486	Sequence 68, Appl
22	208.5	6.8	486	Sequence 72, Appl
23	208.5	6.8	486	Sequence 73, Appl
24	208.5	6.8	502	Sequence 1209, A
25	202.5	6.6	345	Sequence 112, Ap
26	201	6.6	492	Sequence 65, Appl
27	198.5	6.5	484	Sequence 58, Appl

RESULT 1
 US-09-327-984A-38

Sequence 38, Application US/09327984A
 ; Patent No. 6,66854
 ; GENERAL INFORMATION:
 ; APPLICANT: Doetsch, Paul W.
 ; APPLICANT: Kaur, Balveen
 ; APPLICANT: Avery, Angela M.
 ; TITLE OF INVENTION: Broad Specificity DNA Damage Endonuclease
 ; FILE REFERENCE: 25-98
 ; CURRENT APPLICATION NUMBER: US/09/327, 984A
 ; CURRENT FILING DATE: 1999-06-08
 ; PRIOR APPLICATION NUMBER: - US 60,088, 521
 ; PRIOR FILING DATE: 1998-06-08
 ; PRIOR FILING DATE: 1999-05-18
 ; NUMBER OF SEQ ID NOS: 39
 ; SOFTWARE: PatentIn Ver. 2.0
 ; SEQ ID NO: 38
 ; LENGTH: 580
 ; TYPE: PRT
 ; ORGANISM: Homo sapiens
 US-09-327-984A-38

Query Match Score: 100.0%; Pred. No. 2.2e-292; Length 580;
 Best Local Similarity 100.0%; Indels 0; Gaps 0;
 Matches 580; Conservative 0; Mismatches 0;

QY 1 MGTGGLSLSLGDGRGAAPTVSSERLVPDPNDLRKEDVAMELERGEGEDEBOMMIKSE 60
 Db 1 MGTGGLSLSLGDGRGAAPTVSSERLVPDPNDLRKEDVAMELERGEGEDEBOMMIKSE 60
 QY 61 CNPLLOPPIASQFGATAGTECRKSVCVGWERMVKOFLGKTAGRFDVYFSPQIKRS 120
 Db 61 CNPLLOPPIASQFGATAGTECRKSVCVGWERMVKOFLGKTAGRFDVYFSPQIKRS 120

QY 121 KSSLANLYHKGNTTSIPEDEPTVSKRGKSYKCSMAALTSHIQNOSINNSNMLRT 180
 Db 121 KSSLANLYHKGNTTSIPEDEPTVSKRGKSYKCSMAALTSHIQNOSINNSNMLRT 180

QY 181 RSKKKKVFMPPSSSSIQESQGLSNITSTHLLKEEGVDDVNPRURPKKGKVTIKG 240
 Db 181 RSICKKVKVMPSSSSIQESQGLSNITSTHLLKEEGVDDVNPRURPKKGKVTIKG 240

QY 301 SVTSEENSLVKKERSISGSISGFCBKTSGINKRERAKOSEHNIEKYEDTLESETIGT 360
 Db 301 SVTSEENSLVKKERSISGSISGFCBKTSGINKRERAKOSEHNIEKYEDTLESETIGT 360

RESULT 2
US-09-629-222A-2
; Sequence 2, Application US/09629222A
; Patent No. 6599700
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; FILE REFERENCE: FCCC 96 21
; CURRENT APPLICATION NUMBER: US/09/629, 222A
; CURRENT FILING DATE: 2000-07-31
; PRIORITY NUMBER: 09/463, 891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-27
; PRIOR APPLICATION NUMBER: 60/053, 936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FastSEQ For Windows Version 3.0
; LENGTH: 580
; SEQ ID NO: 2
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-629-222A-2

Query Match 100.0%; Score 3055; DB 4; Length 580;
Best Local Similarity 100.0%; Pred. No. 2.e-292;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MGTTGLESLSLGDRGAAPTVTSSERLVPDPNDLKEDVAMELERVGDEBEMMIKRSSE 60
Db 1 MGTTGLESLSLGDRGAAPTVTSSERLVPDPNDLKEDVAMELERVGDEBEMMIKRSSE 60

QY 61 CNPLIQLQPPIASAQFGATAGTCERKSUPCGNBRVVIQRLFKTAGRFDVFIISPQGLKRS 120
Db 61 CNPLIQLQPPIASAQFGATAGTCERKSUPCGNBRVVIQRLFKTAGRFDVFIISPQGLKRS 120

QY 121 KSSLANYLHKNGETSLKPEDFTVLSKRGKRSYKDCSMALTHSLHQNSNNNNWIRT 180
Db 121 KSSLANYLHKNGETSLKPEDFTVLSKRGKRSYKDCSMALTHSLHQNSNNNNWIRT 180

QY 181 RSKCKDVFMPSSSELOESRGLSNFTSTHLLKEDGVDDYFRKVKPKVTLKG 240
Db 181 RSKCKDVFMPSSSELOESRGLSNFTSTHLLKEDGVDDYFRKVKPKVTLKG 240

QY 241 IPIKKTKGKCRKSCSGFVQDSKRSVNCNQDAESEPVQAKSOLDRTVTSAGACGT 300
Db 241 IPIKKTKGKCRKSCSGFVQDSKRSVNCNQDAESEPVQAKSOLDRTVTSAGACGT 300

QY 301 SVTSEENSLVKKERSLSGSNSFCSEQTKGINKFCSAKDSHENEKYEDTELEBEIGT 360
Db 301 SVTSEENSLVKKERSLSGSNSFCSEQTKGINKFCSAKDSHENEKYEDTELEBEIGT 360

QY 361 KVEVERKEHLHTDILKRGSEMDNCSPTRKDFTGKFOEDTPRTQIERRTSYFSS 420
Db 361 KVEVERKEHLHTDILKRGSEMDNCSPTRKDFTGKFOEDTPRTQIERRTSYFSS 420

QY 421 KYNEKALSPPRKAFKPKTTPRSPNLUQETLFHDPMWLIATIFUNTSGKMAIPWK 480
Db 421 KYNEKALSPPRKAFKPKTTPRSPNLUQETLFHDPMWLIATIFUNTSGKMAIPWK 480

QY 421 KYNEKALSPPRKAFKPKTTPRSPNLUQETLFHDPMWLIATIFUNTSGKMAIPWK 480
Db 421 KYNEKALSPPRKAFKPKTTPRSPNLUQETLFHDPMWLIATIFUNTSGKMAIPWK 480

QY 481 FLEKYPSEAERTADWRDVSELLKPLGLYDLRAKTIKVEFSDEYLTKQWKPTELHGIGKY 540
Db 481 FLEKYPSEAERTADWRDVSELLKPLGLYDLRAKTIKVEFSDEYLTKQWKPTELHGIGKY 540

QY 541 GNDSYRICCVENQVHEDHKUNKYHDLWENHEKLISL 580
Db 541 GNDSYRICCVENQVHEDHKUNKYHDLWENHEKLISL 580

RESULT 3
US-09-657-013-56
; Sequence 56, Application US/09657013
; Patent No. 6709817
; GENERAL INFORMATION:
; APPLICANT: Zoghbi, Huda Y.
; APPLICANT: Van den Veyver, Ignatia B
; APPLICANT: Amir, Ruthie
; APPLICANT: Francke, Uta
; TITLE OF INVENTION: Methods of Identifying Mutations in a Methyl-CPG-Binding Domain
; FILE REFERENCE: HO-20183US1/09905371
; CURRENT APPLICATION NUMBER: US/09/657, 013
; CURRENT FILING DATE: 2000-03-07
; PRIORITY NUMBER: US 60/152, 778
; PRIOR FILING DATE: 1999-09-07
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO: 56
; LENGTH: 580
; TYPE: PRT
; ORGANISM: Human
; US-09-657-013-56

Query Match 100.0%; Score 3055; DB 4; Length 580;
Best Local Similarity 100.0%; Pred. No. 2.e-292;
Matches 580; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MGTTGLESLSLGDRGAAPTVTSSERLVPDPNDLKEDVAMELERVGDEBEMMIKRSSE 60
Db 1 MGTTGLESLSLGDRGAAPTVTSSERLVPDPNDLKEDVAMELERVGDEBEMMIKRSSE 60

QY 61 CNPLIQLQPPIASAQFGATAGTCERKSUPCGNBRVVIQRLFKTAGRFDVFIISPQGLKRS 120
Db 61 CNPLIQLQPPIASAQFGATAGTCERKSUPCGNBRVVIQRLFKTAGRFDVFIISPQGLKRS 120

QY 121 KSSLANYLHKNGETSLKPEDFTVLSKRGKRSYKDCSMALTHSLHQNSNNNNWIRT 180
Db 121 KSSLANYLHKNGETSLKPEDFTVLSKRGKRSYKDCSMALTHSLHQNSNNNNWIRT 180

QY 181 RSKCKDVFMPSSSELOESRGLSNFTSTHLLKEDGVDDYFRKVKPKVTLKG 240
Db 181 RSKCKDVFMPSSSELOESRGLSNFTSTHLLKEDGVDDYFRKVKPKVTLKG 240

QY 241 IPIKKTKGKCRKSCSGFVQDSKRSVNCNQDAESEPVQAKSOLDRTVTSAGACGT 300
Db 241 IPIKKTKGKCRKSCSGFVQDSKRSVNCNQDAESEPVQAKSOLDRTVTSAGACGT 300

QY 301 SVTSEENSLVKKERSLSGSNSFCSEQTKGINKFCSAKDSHENEKYEDTELEBEIGT 360
Db 301 SVTSEENSLVKKERSLSGSNSFCSEQTKGINKFCSAKDSHENEKYEDTELEBEIGT 360

QY 361 KVEVERKEHLHTDILKRGSEMDNCSPTRKDFTGKFOEDTPRTQIERRTSYFSS 420
Db 361 KVEVERKEHLHTDILKRGSEMDNCSPTRKDFTGKFOEDTPRTQIERRTSYFSS 420

QY 421 KYNEKALSPPRKAFKPKTTPRSPNLUQETLFHDPMWLIATIFUNTSGKMAIPWK 480
Db 421 KYNEKALSPPRKAFKPKTTPRSPNLUQETLFHDPMWLIATIFUNTSGKMAIPWK 480

RESULT 4
US-09-629-222A-24
; Sequence 24, Application US/0962922A
; Patent No. 659700
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/09/629, 222A
; CURRENT FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463, 891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053, 936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 24
; LENGTH: 574
; ORGANISM: Homo sapiens
US-09-629-222A-24
Query Match 98.6%; Score 3011; DB 4; Length 574;
Best Local Similarity 99.0%; Pred. No. 4,7e-288;
Matches 574; Conservative 0; Mismatches 0; Indels 6; Gaps 1;
QY |||||MGITGLLSLIGRGAAPTVTSSERLVDPNDLKEVAMELSERDEEPMKRSSE 60
Db 1 MGTGLESLSLGDRGAAPTVTSSERLVDPNDLKEVAMELSERDEEPMKRSSE 60
QY |||||CMLLQBETIASAGTAGATGTECKSVPGCWGRWRYVKORLFGKTYAGRDFVYFSPQGLKFRS 120
Db 61 CMLLQBETIASAGTAGATGTECKSVPGCWGRWRYVKORLFGKTYAGRDFVYFSPQGLKFRS 120
QY |||||CNPLOQEPITASAOFGATGTECKSVPGCWGRWRYVKORLFGKTYAGRDFVYFSPQGLKFRS 120
Db 61 CNPLOQEPITASAOFGATGTECKSVPGCWGRWRYVKORLFGKTYAGRDFVYFSPQGLKFRS 120
QY |||||KSSLANYLHKGNTSLKEBFDFTVLSGRKGSRYKOCMSMALLSHLQNQNSNNMLRT 180
Db 121 KSSLANYLHKGNTSLKEBFDFTVLSGRKGSRYKOCMSMALLSHLQNQNSNNMLRT 180
QY |||||181 RSKCKKDVFMPASSSEIQLBSRGISNFSTHLALKEDGVDDNFRKPKKVTLKG 240
Db 181 RSKCKKDVFMPASSSEIQLBSRGISNFSTHLALKEDGVDDNFRKPKKVTLKG 240
QY |||||241 IPIKTKKGCRKCSGFTQDSKRESVENKADESEPYAQKSOLDRTVCISDAGCETL 300
Db 241 IPIKTKKGCRKCSGFTQDSKRESVENKADESEPYAQKSOLDRTVCISDAGCETL 300
QY |||||301 SVTSEENSLVKKERSLSSGSNCSEQTKGINKFCSAKDSHENEKYEDTELESEEGT 360
Db 301 SVTSEENSLVKKERSLSSGSNCSEQTKGINKFCSAKDSHENEKYEDTELESEEGT 360
QY |||||361 KVEVERKEHLHTDLKRSSEMDNCSPTRKOTGEKFKQDQIPQRERRTSIYESS 420
Db 361 KVEVERKEHLHTDLKRSSEMDNCSPTRKOTGEKFKQDQIPQRERRTSIYESS 420
QY |||||421 KYNEALSPRRKPKOMPSPRLNYVQETLFDWKLLIATFLNRUTSGRMAIPVWAK 480
Db 421 KYNEALSPRRKPKOMPSPRLNYVQETLFDWKLLIATFLNRUTSGRMAIPVWAK 480
QY |||||415 KYNEALSPRRKPKOMPSPRLNYVQETLFDWKLLIATFLNRUTSGRMAIPVWAK 474
Db 415 KYNEALSPRRKPKOMPSPRLNYVQETLFDWKLLIATFLNRUTSGRMAIPVWAK 474

RESULT 5
US-09-629-222A-29
; Sequence 29, Application US/0962922A
; Patent No. 659700
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/09/629, 222A
; CURRENT FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463, 891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053, 936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 29
; LENGTH: 384
; TYPE: PRT
; ORGANISM: Mus musculus
US-09-629-222A-29
Query Match 43.4%; Score 1326; DB 4; Length 384;
Best Local Similarity 54.3%; Pred. No. 5,3e-122; Indels 138; Gaps 4;
Matches 283; Conservative 29; Mismatches 71; Indels 138; Gaps 4;
QY |||||36 KEDVAMELERVEDEBCOMMIKRSSECNPLORPIASAQFGATAGTCRKSVP CGWERYVK 95
Db 1 KEDIAVGIGGYCGBDKLIVI--SSERSLJOOPTAST-LSSRATBQHKPVPCGWERVK 57
QY |||||96 ORLFGKTAGRFDVYFSPQGLKFRS 155
Db 58 ORLSGKTAGKFVYFSPQGLKFRS 117
QY |||||156 KOCSMALLSHLQNQNSNNMLRTSKCKDQVMPSSSBLOESGLSNFTSHLKL 215
Db 118 KHOSLAAALTSLOPNETDVSKONIKTRSKWKTDLVPLPLSGTISPSSGGLSNSNACILUR 177
QY |||||216 EDBGVDVNFRCRKPKKVTLKGTPRIKTKKGCRKCSGFTQDSKRESVENKADES 275
Db 178 EHHDIOVDESEKRKSKRKVTLKGTAQTKQKCRKSLLESTORURKRA----- 228
QY |||||276 EPVAQKSQOLDRTVCISDAGCETLSTVSEENSLVKKERSLSSGSNCSEQTKG 335
Db 229 -----
QY |||||336 FCGSAKDSHENEKYEDTFLSERBIGTKVVERKEHLHTDLKRSSEMDNCSPTRKPKFTG 395
Db 229 -----
QY |||||396 BKIQFQDTPTRQERKTSIYFSSKNCSEQTKGINKFCSAKDSHENEKYEDTELESEEGT 455
Db 229 -----
QY |||||456 PWKLLIATFLNRUTSGRMAIPVWAKFQEVARTADWRDVSEBLKPIGLYDRAKT 515
Db 284 PWKLLIATFLNRUTSGRMAIPVWAKFQEVARTADWRDVSEBLKPIGLYDRAKT 543
QY |||||516 TPKSDDELTOKWPKYBLHGIG-KYNSDYSRIFCVNEWKO 555
Db 344 IKESDELTOKWPKYBLHGIGWIKYONDYSRIFCVNEWKO 384


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; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 32
; LENGTH: 126
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-629-222A-32

Query Match 22.8%; Score 697; DB 4; Length 126;
Best Local Similarity 100.0%; Pred. No. 1e-60; Matches 126; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Prior Application Number: 60/053,936
Prior Filing Date: 1997-07-28
Number of SEQ ID NOS: 73
Software: FastSEQ for Windows Version 3.0
SEQ ID NO 37
Length: 85
Type: PRT
Organism: Homo sapiens
US-09-629-222A-37

Query Match 14.5%; Score 443; DB 4; Length 85;
Best Local Similarity 100.0%; Pred. No. 6.3e-36; Matches 85; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Prior Application Number: 60/048,891
Prior Filing Date: 1997-07-28
Number of SEQ ID NOS: 73
Software: FastSEQ for Windows Version 3.0
SEQ ID NO 30
Length: 119
Type: PRT
Organism: Homo sapiens
US-09-629-222A-30

Query Match 20.2%; Score 616; DB 4; Length 119;
Best Local Similarity 100.0%; Pred. No. 9.3e-53; Matches 119; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Prior Application Number: US/09/629,222A
Prior Filing Date: 2000-01-28
Prior Application Number: PCT/US98/15828
Prior Application Number: 60/053,936
Prior Filing Date: 1997-07-28
Number of SEQ ID NOS: 73
Software: FastSEQ for Windows Version 3.0
SEQ ID NO 30
Length: 119
Type: PRT
Organism: Homo sapiens
US-09-629-222A-30

; RESULT 8
; Sequence 30, Application US/09629222A
; Patent No. 6599700
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/09/629,222A
; CURRENT FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-07-28
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 30
; LENGTH: 119
; TYPE: PRT
; ORGANISM: Homo sapiens

; RESULT 9
; Sequence 37, Application US/09629222A
; Patent No. 6599700
; GENERAL INFORMATION:
; APPLICANT: Bellacosa, Alfonso
; TITLE OF INVENTION: Methods for Detection of Transition
; TITLE OF INVENTION: Single-Nucleotide Polymorphisms
; FILE REFERENCE: FCCC 96-21
; CURRENT APPLICATION NUMBER: US/09/629,222A
; CURRENT FILING DATE: 2000-07-31
; PRIOR APPLICATION NUMBER: 09/463,891
; PRIOR FILING DATE: 2000-01-28
; PRIOR APPLICATION NUMBER: PCT/US98/15828
; PRIOR FILING DATE: 1998-07-28
; PRIOR APPLICATION NUMBER: 60/053,936
; PRIOR FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,891
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,020
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,876
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,895
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,884
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,894
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,971
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,882
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,899
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,893
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,900
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,901

; RESULT 10
; Sequence 1100, Application US/09205258
; Patent No. 6525174
; GENERAL INFORMATION:
; APPLICANT: Young et al.
; TITLE OF INVENTION: 207 Human Secreted Proteins
; FILE REFERENCE: P2007P1
; CURRENT APPLICATION NUMBER: US/09/205,258
; CURRENT FILING DATE: 1998-12-04
; EARLIER APPLICATION NUMBER: PCT/US98/11422
; EARLIER FILING DATE: 1998-06-04
; EARLIER APPLICATION NUMBER: 60/048,885
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,375
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,881
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,880
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,896
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/049,020
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; EARLIER APPLICATION NUMBER: 60/048,876
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,895
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,884
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,894
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,971
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,882
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,899
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,893
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,900
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/048,901

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EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,892
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,915
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/049,019
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,970
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,972
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,916
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/049,373
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,875
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/049,374
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,917
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,949
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,974
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,883
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,897
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,963
 EARLIER FILING DATE: 1997-06-06
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 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,962
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,963
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,977
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/048,878
 EARLIER FILING DATE: 1997-06-06
 EARLIER APPLICATION NUMBER: 60/070,923
 EARLIER FILING DATE: 1997-12-18
 EARLIER APPLICATION NUMBER: 60/092,921
 EARLIER FILING DATE: 1998-07-15
 EARLIER APPLICATION NUMBER: 60/094,657
 NUMBER OF SEQ ID NOS: 1227
 SOFTWARE: PatentIn Ver. 2.0
 SEQ ID NO: 1100
 LENGTH: 50
 TYPE: PRT
 ORGANISM: Homo sapiens

RESULT 11
 US-09-657-013-69
 ; Sequence 69, Application US/09657013
 ; Patent No. 6709817
 GENERAL INFORMATION:
 ; APPLICANT: Zoghlbi, Huda Y.
 ; APPLICANT: Van den Veyver, Ignatia B
 ; APPLICANT: Amir, Ruthie
 ; APPLICANT: Francke, Uta
 TITLE OF INVENTION: Methods of Identifying Mutations in a Methyl-CPG-Binding Domain
 TITLE OF INVENTION: Methods of Identifying Mutations in a Methyl-CPG-Binding Domain and Tre
 FILE REFERENCE: HO-01893US1/09905371
 CURRENT APPLICATION NUMBER: US/09/657,013
 CURRENT FILING DATE: 2000-09-07
 PRIOR APPLICATION NUMBER: US 60/152,773
 PRIOR FILING DATE: 1999-09-07
 NUMBER OF SEQ ID NOS: 114
 SOFTWARE: PatentIn version 3.1
 SEQ ID NO: 70
 LENGTH: 467
 TYPE: PRT
 ORGANISM: Frog
 US-09-657-013-69
 ;
 Query Match 8.3%: Score 254; DB 4; Length 50;
 Best Local Similarity 98.0%; Pred. No. 1.2e-17; Mismatches 1; Indels 0; Gaps 0;
 Matches 49; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 110 FSSPOGIKFRKSLSLANYLNHNGETSIKPERPDTFLSKRGKIKSRKDGS 159
 Db 1 FSSPOGIKFRKSLSLANYLNHNGETSIKPERPDTFLSKRGKIKSRKDGS 50
 ;
 RESULT 12
 US-09-657-013-70
 Sequence 70, Application US/09657013
 PATENT NO. 6709817
 GENERAL INFORMATION:
 APPLICANT: Zoghlbi, Huda Y.
 APPLICANT: Van den Veyver, Ignatia B
 APPLICANT: Amir, Ruthie
 APPLICANT: Francke, Uta
 TITLE OF INVENTION: Methods of Identifying Mutations in a Methyl-CPG-Binding Domain and Tre
 FILE REFERENCE: HO-01893US1/09905371
 CURRENT APPLICATION NUMBER: US/09/657,013
 CURRENT FILING DATE: 2000-09-07
 PRIOR APPLICATION NUMBER: US 60/152,773
 PRIOR FILING DATE: 1999-09-07
 NUMBER OF SEQ ID NOS: 114
 SOFTWARE: PatentIn version 3.1
 SEQ ID NO: 70
 LENGTH: 467
 TYPE: PRT
 ORGANISM: Frog
 US-09-657-013-70
 ;
 Query Match 7.2%: Score 220.5; DB 4; Length 467;
 Best Local Similarity 25.7%; Pred. No. 1e-12; Mismatches 157; Indels 67; Gaps 15;

QY 22 SSERLVLVPDPMDPLRKEDVAMELER--VGEDEBQMMIKRS--SECNPILQEPIASAQFGA 76
 Db 49 SSEH-QPGERPAPDEGKADMSSAENLAUVPASSASPRQRSPVTRDRGPMPYEDP----- 99

QY 77 TAGTECRKSVCVGWERRVKORLGKTAGRFVYFISPOGLKFRSKSSLANY 136
 Db 100 -----TLPBGWTRKIKQRKGSRSGAKFEDVYLINPNQKAFRSKVELIAYFQKGDTSL 151

QY 137 KPEDFDPFTVLSKRGTSRKYKDCSMAALTSHLQNQNSNNWNLRTSKCKD 187
 Db 152 DPDNDFDFTV-TGRGSPSR-----KOPKKPKRSSVSGGRGRPKGSKKVPPVK 204

QY 195 SSBLOBSRGLSNFTSHLLK----EDEGTDVNWRKV----RKPCKVTLKGIP 242
 Db 205 SEGVQVKRVTK-SPEKLVLVNPYSGTKEADATTSQVNLVTKRGCRKRE-TDPSAAP 262

QY 243 IRTKKGCRKCSGFVQSDSRESVKADESEPAQSKOLDRTVCISDAGCETLSV 302
 Db 263 KGRGRKPSNVSLAAAABAAKKAI--KESSIKPLIE---TULPIKGKRKTISV 313

QY 303 TSEE-----NSLVKK-----KERLSSSNFCSEOKTSGIINFKCSADSENEKYE 349
 Db 314 DVKDTRPEPITPVIEKUMQJONPAKSPESRSTEGSPKIKIGLPRKELQOHHHHHH 373

QY 350 DTFLESBEGTKVEVERKEIL 371
 Db 374 HHHSSEKASATSPPEPTSKDN 395

RESULT 13
 US-09-657-013-74
 ; Sequence 74, Application US/09657013
 ; Patent No. 6709817
 ; GENERAL INFORMATION:
 ; APPLICANT: Zoghbi, Huda Y.
 ; INVENTION: Methods of Identifying Mutations in a Methyl-CPG-Binding Domain and Tre
 ; APPLICANT: Van den Veyver, Ignatia B
 ; APPLICANT: Amir, Rutchie
 ; APPLICANT: Francke, Utta
 ; TITLE OF INVENTION: Methods of Identifying Mutations in a Methyl-CPG-Binding Domain and Tre
 ; FILE REFERENCE: HO-P01893US1/09905371
 ; CURRENT APPLICATION NUMBER: US/09/657,013
 ; CURRENT FILING DATE: 2000-09-07
 ; PRIORITY NUMBER: US 60/152,778
 ; PRIORITY FILING DATE: 1999-09-07
 ; NUMBER OF SEQ ID NOS: 114
 ; SOFTWARE: Patentin version 3.1
 ; SEQ ID NO: 75
 ; LENGTH: 476
 ; TYPE: PRT
 ; ORGANISM: Human
 ; US-09-657-013-75

Query Match 6.9%; Score 209; DB 4; Length 476;
 Best Local Similarity 24.0%; Pred. No. 1.4e-11;
 Matches 104; Conservative 53; Mismatches 149; Indels 128; Gaps 17;

QY 37 EDVAMELERGEBEQQMIRKSECBNPILQEPIASAQFG-----ATAGT 81
 Db 13 KDKPLKKVKKKBKEKEKGHEPVQPSAHHSAAEPAEAGRAETSBGGSARLCEASAASP 72
 QY 82 CRKSV-----PGCWERVVKORLGKTAGRFVYFISPOGLKFRSKSSLANYL 128
 Db 73 ORRSIIIRDGPWMYDDPPTLPGCWTRKIKQRKGSRSGAKYDVLINPQKAFRSKVELIAYF 132

QY 129 HKNGETSLKEPDFFTVLSKRGTSRKYKDCSMAALTSHLQNQNSNNWNLRTSKCKD 188
 Db 133 EKVGDTSLDPDFTV-TGRGSPSR-----REQPKK- 165

QY 189 FMPSSSELQESRGL--SNFTSHLLKEDEGYDUDVNFRKVRKPKGVTLKGIP 245
 Db 166 ---PKSFKAPGPGRGRPKGSGITRPKAATSEGQVK--RVLERSPK--LLVQMPF-Q 217

QY 246 TKGGCRKSCGFVQSPS---KRESYVKADAESPVAKSOLDRTVCISDAGCETLSV 301
 Db 218 TSPGKGKEGGATTSTQVMVTKRPGRKKAADPOQKPKR-----GRKGGSVA 267

QY 302 VTSEBNLSVKKKERSLSSGSNCSEOKTSGIINKCSAKOSENHEKEDTFLESIGTK 361
 Db 268 AAAAKKKKAVKKESSR-----SYQETVLPK-----RKTRET-----VS 303

QY 362 VEVVERKEHLHTDL--KRESEMDDNNCSPTRDFTGKIFQBDTIPRTOTERRKTSLYFS 419
 Db 304 IEVKVKPLLVSTLGKSGKGLKTCSPRK-----SKESSPKGR----- 344

QY 245 KTKKGCRKSCSGFVQSDS---KRESYVKADAESPVAKSOLDRTVCISDAGCETLSV 300
 Db 227 QTPSPGKACBEGGATTSTQVMVTKRPGRKKAADPOQKPKR-----GRKGGSV 276

QY 301 SVTSBENSLVKKKERSLSSGSNCSEOKTSGIINKCSAKOSENHEKEDTFLESIGTK 360
 Db 277 AAAABE---AKKK---AVKGSSSTRSVOETVLPK-----RKTRET-----V 312

QY 361 KVEVVERKEHLHTDL--KRGSEMDNNCSPTRKDFTGKIFQBDTIPRTQIERKTSLYF 418
 Db 313 SIEVEVKVLLVSLIGERSGKGKLTCKSPRK-----SKESSPKGR----- 354

QY 419 SSXKVKRKAASPPRK 433
 Db 355 ---SSASSPPK 365

RESULT 15
 US-09-657-013-63
 ; Sequence 63 Application US/09657013
 ; Patent No. 6709817
 ; GENERAL INFORMATION:
 ; APPLICANT: Zoghbi, Huda Y.
 ; APPLICANT: van den Veyver, Ignatia B
 ; APPLICANT: Amir, Ruthie
 ; APPLICANT: Francke, Uta
 TITLE OF INVENTION: Methods of Identifying Mutations in a Methyl-CPG-Binding Domain
 FILE NUMBER: HO-P01893US1/09905371
 CURRENT APPLICATION NUMBER: US/09/657,013
 CURRENT FILING DATE: 2000-09-07
 PRIOR APPLICATION NUMBER: US 60/152,778
 PRIOR FILING DATE: 1999-09-07
 NUMBER OF SEQ ID NOS: 114
 SOFTWARE: PatentIn version 3.1
 SEQ ID NO 63
 LENGTH: 477
 TYPE: PRT
 ORGANISM: Human
 US-09-657-013-63

Query Match 6.8%; Score 208.5; DB 4; Length 477;
 Best Local Similarity 23.9%; Preq. No. 1.6e-11; Matches 149; Mismatches 149; Gaps 17;
 Matches 104; Conservative 53; Indels 129;

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QY      37  EDVAMELER/CBDEBOMMIKSECNPLLOPIASAQFG-----ATAGT 80
       ; | : :| :||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:|
       13 KDKPKUFKKVKDKKEKEKGHEPVQPSAHSABPAGGETSSGSGSAPAVPEASAP 72
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:|
       81 ECRKSV-----PGWERYVKQRLFGKTPAGRFDWYFSPQGLKFRSKSLANY 127
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:|
       73 FQRRSIIRDQPGMYDPTLPGWTRKIKQKGRSGRSAGKYDWYLINPOGKAFRSKVELIAY 132
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:|
       128 LHKNGATSLIKEPEDOPTVLSKRGTSRKYKOCSSMAALTSHQINQNSNNMLRTSKCKD 187
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:|
       133 FEKVGDTSLDNDPDPYV-TGKGSPSR-----REOKPPK- 166
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:|
       188 VFMPSSSSELQESGL---SNFTSHLLIKEDEGVDDYAFFRKPKPKGVTLKGIPK 244
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:|
       167 ---BKSPKAPGTGGRGRPGKSGCTRPKATSECWQVK--RVLKSPGK--LVTKMPF- 217
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:|
       245 RTKKGCRKSCSGFVOSDS---KRESVCNKDAESEPVAQKSQDRTVCISDAGACETL 300
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:|
       218 OTSPGGKAEGGGATTSTQVMVIKRPFORKKBAFDQAPKRR-----GRKPGSV 267
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:|
       301 SVTSSSENLVKKKERSLSSSNCBQKSGGIINIFCSAKDSENEKEDTFLESEEIGT 360
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:||:||:||:||:|
       268 AAAAAEAKKKAVKESSIR-----SVOETVLPKK-----RKTRBT-----V 303
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:||:||:||:|
       361 KVEVVERKEHHTDIL---KQGSEMDNCSSTRKDFTGEKIFFQEDTIPIQIERKTSLYF 418
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:||:||:||:|
       304 SIEVKWKVQLLVLSTIGEKSGKGLKTCSPGRK-----SKESSPGR----- 345
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:||:|
       QY      419 SKYKNEALSPPRK 433
       ; | :|:||:||:||:||:||:||:||:||:||:||:||:|
       Db      346 ---SSSASSPPKKE 356
  
```